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(71) Sökande AstraZeneca AB, Södertälje SE
Applicant (s)

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THERAPEUTIC AGENTS

Field of the invention

5 The present invention relates to certain novel benzoic acid derivatives, to processes for preparing such compounds, to their utility in treating clinical conditions associated with insulin resistance, to methods for their therapeutic use and to pharmaceutical compositions containing them.

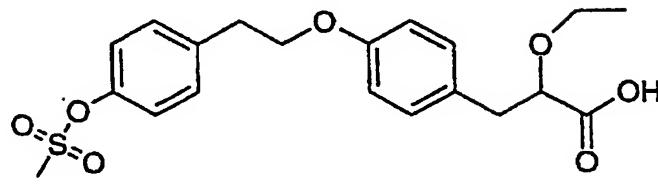
Background of the invention

10 The Insulin Resistance Syndrome (IRS) including type 2 diabetes mellitus, which refers to a cluster of manifestations including insulin resistance with accompanying hyperinsulinaemia, possible type 2 diabetes mellitus, arterial hypertension, central 15 (visceral) obesity, dyslipidaemia observed as deranged lipoprotein levels typically characterised by elevated VLDL (very low density lipoproteins), small dense LDL particles and reduced HDL (high density lipoprotein) concentrations and reduced fibrinolysis.

20 Recent epidemiological research has documented that individuals with insulin resistance run a greatly increased risk of cardiovascular morbidity and mortality, notably suffering from myocardial infarction and stroke. In type 2 diabetes mellitus atherosclerosis related conditions cause up to 80% of all deaths.

25 In clinical medicine there is awareness of the need to increase the insulin sensitivity in IRS suffering patients and thus to correct the dyslipidaemia which is considered to cause the accelerated progress of atherosclerosis. However, currently this is not a universally well defined disease.

The S-enantiomer of the compound of formula C below



C

2-ethoxy-3-[4-(2-(4-methanesulfonyloxyphenyl)ethoxy)phenyl]propanoic acid, is disclosed in PCT Publication Number WO99/62872. This compound is reported to be a modulator of peroxisome proliferator-activated receptors (PPAR, for a review of the PPARs see T. M. Willson et al, J Med Chem 2000, Vol 43, 527) and has combined PPAR α /PPAR γ agonist activity (Structure, 2001, Vol 9, 699, P. Cronet et al). This compound is effective in treating conditions associated with insulin resistance.

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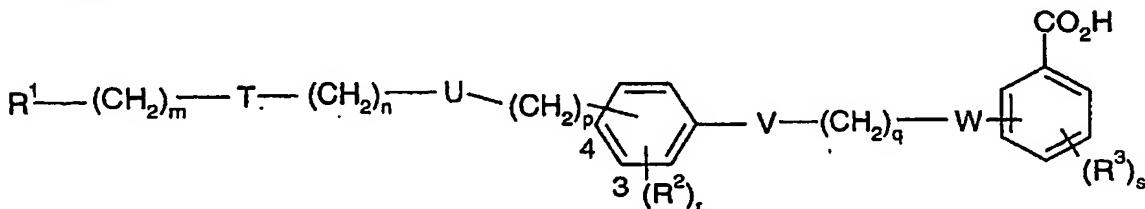
Surprisingly a series of compounds has now been found which are modulators of PPAR α and/or PPAR γ activity.

Description of the invention

15

The present invention provides a compound of formula I

A compound of formula I



20

wherein

R^1 represents aryl optionally substituted by a heterocyclic group or a heterocyclic group optionally substituted by aryl wherein each aryl or heterocyclic group is optionally substituted by one or more of the following groups:

- a C_{1-6} alkyl group;
- 5 a C_{1-6} acyl group;
aryl C_{1-6} alkyl, wherein the alkyl, aryl, or alkylaryl group is optionally substituted by one or more R^b ;
- halogen,
- CN and NO_2 ,
- 10 - NR^cCOOR^a ;
- NR^cCOR^a ;
- NR^cR^a ;
- $NR^cSO_2R^d$;
- $NR^cCONR^kR^c$;
- 15 - $NR^cCSNR^aR^k$;
- OR^a ;
- OSO_2R^d ;
- SO_2R^d ;
- SOR^d ;
- 20 - SR^c ;
- $SO_2NR^aR^f$;
- SO_2OR^a ;
- $CONR^cR^a$;
- $OCONR^fR^a$;
- 25 wherein R^a represents H, a C_{1-6} alkyl group, aryl or aryl C_{1-6} alkyl group wherein the alkyl, aryl or aryl C_{1-6} alkyl group is optionally substituted one or more times by R^b , wherein R^b represents C_{1-6} alkyl, aryl, aryl C_{1-6} alkyl, cyano, - NR^cR^d , =O, halo, -OH, -SH, - OC_{1-4} alkyl, -Oaryl, - OC_{1-4} alkylaryl, - COR^c , - SR^d , - SOR^d , or - SO_2R^d , wherein R^c represents H, C_{1-4} alkyl, aryl, aryl C_{1-4} alkyl and R^d represents C_{1-4} alkyl, aryl, aryl C_{1-4} alkyl;
- 30 wherein R^f represents hydrogen, C_{1-4} alkyl, C_{1-4} acyl, aryl, aryl C_{1-4} alkyl and R^a is as defined above; and
 R^k represents hydrogen, C_{1-4} alkyl, aryl, aryl C_{1-4} alkyl;

m represents 0, 1, 2, 3, 4 or 5 ;

5 T represents O, S, NC(O)N(R⁴), S(O₂)N(R⁵), (R⁵)NS(O₂), N(R⁶)C(O), C(O)N(R⁷), or a single bond;

n represents 0, 1, 2, 3, 4 or 5 ;

10 U represents O, S or a single bond provided that when U is O or S then n represents 1, 2, 3, 4 or 5 and further provided that T and U do not both represent a single bond simultaneously;

p represents 0, 1, 2, 3, 4 or 5 ;

15 wherein the group (CH₂)_p, or the group U if p is 0, is attached at the 3 or 4 position of the phenyl ring as indicated in formula I

V represents O, S, NR⁸, or a single bond;

20 q represents 1, 2 or 3 ;

W represents O, S, N(R⁹)C(O) , NR¹⁰, or a single bond;

25 R² represents halo, a C₁₋₄alkyl group which is optionally substituted by one or more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro, a C₁₋₄acyl group, aryl, an arylC₁₋₄alkyl group, CN or NO₂ ;

r represents 0, 1, 2 or 3 ;

30 R³ represents represents halo, a C₁₋₄alkyl group which is optionally substituted by one or more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro, a C₁₋₄acyl group, aryl, an arylC₁₋₄alkyl group, or CN ;

s represents 0, 1, 2 or 3 ; and

R⁴, R⁵, R⁶, R⁷, R⁸, R⁹ and R¹⁰ independently represent H, a C₁₋₁₀alkyl group, aryl or an arylC₁₋₄alkyl group or when m is 0 and T represents a group N(R⁶)C(O) or a group (R⁵)NS(O₂) then R¹ and R⁶ or R¹ and R⁵ together with the nitrogen atom to which they are attached represent a heteroaryl group; as well as optical isomers and racemates thereof as well as pharmaceutically acceptable salts, prodrugs, solvates and crystalline forms thereof.

10

Examples of C₁₋₆alkyl include methyl, ethyl, n-propyl, isopropyl, n-butyl, iso-butyl, sec-butyl, t-butyl and straight- and branched-chain pentyl and hexyl as well as cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl. Preferred alkyl groups are methyl, ethyl, propyl, isopropyl and tertiary butyl.

15 Unless otherwise stated or indicated, the term "halogen" shall mean fluorine, chlorine, bromine or iodine, preferably fluorine.

Unless otherwise stated or indicated, the term "aryl" denotes a substituted or unsubstituted phenyl or a fused ring system such as naphthyl.

20 Unless otherwise stated or indicated, the term "a heterocyclic group" is a saturated, partially saturated or unsaturated, mono or bicyclic ring containing 4-12 atoms of which at least one atom is chosen from nitrogen, sulphur or oxygen, which may, unless otherwise specified, be carbon or nitrogen linked, wherein a -CH₂- group can optionally be replaced by a -C(O)- and a ring sulphur atom may be optionally oxidised to form the S-oxide(s).

25 Examples and suitable values of the term "heterocyclic group" are morpholino, piperidyl, pyridyl, pyranyl, pyrrolyl, imidazolyl, thiazolyl, indolyl, quinolyl, isoquinolyl, thienyl, 1,3-benzodioxolyl, 1,3-dioxolanyl, thiadiazolyl, piperazinyl, isothiazolidinyl, 1,3,4-triazolyl, tetrazolyl, pyrrolidinyl, 2-oxazolidinonyl, 5-isoxazolonyl, benz-3-azepinyl, 1,4-benzodioxanyl, thiomorpholino, pyrrolinyl, homopiperazinyl, 3,5-dioxapiperidinyl, 3-pyrazolin-5-onyl, tetrahydropyrananyl, benzimidazolyl, benzthiazolyl,

30 imidazo[1,2-a]pyridyl, pyrimidyl, pyrazinyl, pyridazinyl, isoxazolyl, 4-pyridone, 1-isoquinolone, 2-pyrrolidone, 4-thiazolidone, dihydroisoquinol-2(1H)-yl, 2,3-dihydro-1,5-benzothiazepin-4(5H)-one. Preferably a "heterocyclic group" is pyridyl,

imidazolyl, thiazolyl, quinolyl, thienyl, 1,3-benzodioxolyl, 1,3-dioxolanyl, isothiazolidinyl, 1,3,4-triazolyl, tetrazolyl, 2-oxazolidinonyl, 5-isoxazolonyl, benz-3-azepinyl, hydantoinyl, 1,4-benzodioxanyl, thiomorpholino, 3-pyrazolin-5-onyl, benzimidazolyl, benzthiazolyl, imidazo[1,2-a]pyridyl, pyrimidyl, pyrazinyl, and 2,3-dihydro-1,5-benzothiazepin-4(5H)-one.

In another aspect the invention provides a compound of formula I as described immediately above with the provisos that

- 1) when R¹ is phenyl optionally substituted by one or two groups independently selected from halo, a C₁₋₄alkyl group which is optionally substituted by one or more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro;
m is 1;
T is N(R⁶)C(O) wherein R⁶ represents a C₂₋₈alkyl group which is optionally interrupted by oxygen;
- 15 n is 1;
U is absent or represents methylene;
p is 0;
r is 0;
V is O or S;
- 20 q is 1; and
W is a single bond attached to the position ortho to the carboxylic acid group;
then s does not represent 0; and
- 25 2) when R¹ is phenyl optionally substituted by one or two groups independently selected from halo, a C₁₋₄alkyl group which is optionally substituted by one or more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro;
m is 1;
T is N(R⁶)C(O) wherein R⁶ represents an unbranched C₂₋₇alkyl group;
- 30 n is 1;
U is O;
p is 0;
r is 0 or 1;

and when r is 1 R² is attached at the 3 position and is OCH₃;

V is a single bond;

q is 2; and

W is O or S attached to the position ortho to the carboxylic acid group;

5 then s does not represent 0.

Further values of R¹, T, U, V, W, R², R³, m, n, p, q, r and s in compounds of Formula I now follow. It will be understood that such values may be used with any of the definitions, claims or embodiments defined hereinbefore or hereinafter.

10

In a first aspect R¹ represents phenyl which is optionally substituted by one or more of the following: halo, hydroxy, a C₁₋₄alkyl group which is optionally substituted by one or more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro, benzyloxy, a C₁₋₄alkylsulphonyloxy group, phenyl or a heteroaryl group, or R¹ represents a heterocyclic group which is optionally substituted by one or more of the following: halo, a C₁₋₄alkyl group which is optionally substituted by one or more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro or phenyl optionally substituted by one or more of the following: halo, a C₁₋₄alkyl group which is optionally substituted by one or more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro. In particular R¹ represents phenyl, furyl, pyridyl or thiazolyl each of which is optionally substituted by one or more of the following: halo (particularly fluoro), a C₁₋₄alkyl group, trifluoromethyl, a C₁₋₄alkoxy group, methanesulphonyloxy, hydroxy, benzyloxy, imidazolyl or phenyl.

25

In a second aspect the group -(CH₂)_m-T-(CH₂)_n-U-(CH₂)_p- is attached at either the 3 or 4 position in the phenyl ring as indicated by the numbers in formula I and represents a group selected from one or more of the following: O(CH₂)₂, O(CH₂)₃, NC(O)NR⁴(CH₂)₂, CH₂S(O₂)NR⁵(CH₂)₂, CH₂N(R⁶)C(O)CH₂, (CH₂)₂N(R⁶)C(O)(CH₂)₂, C(O)NR⁷CH₂, C(O)NR⁷(CH₂)₂, and CH₂N(R⁶)C(O)CH₂O, wherein R⁴, R⁵, R⁶, and R⁷ are as previously defined.

In a third aspect the group $-(CH_2)_m-T-(CH_2)_n-U-(CH_2)_p-$ is attached at the 4 position in the phenyl ring as indicated by the numbers in formula I, that is para to the group V.

In a fourth aspect the group $-V-(CH_2)_q-W-$ represents a group selected from one or more of the following: OCH_2 , SCH_2 , $NHCH_2$, CH_2CH_2S or CH_2CH_2O .

In a fifth aspect the group $-V-(CH_2)_q-W-$ represents the group OCH_2 .

In a sixth aspect the group $-V-(CH_2)_q-W-$ is joined at the ortho position with respect to the carboxylic acid group.

In a seventh aspect R^2 is halo, a C_{1-4} alkyl group or a C_{1-4} alkoxy group and r is 0 or 1.

It will be appreciated by those skilled in the art that certain compounds of formula I contain an optically active centre and therefore can exist as enantiomers which can be separated as described later. It is expected that most, if not all, of the activity of the compounds of formula I resides in one enantiomer: either the S or the R enantiomer or the (+) or the (-) enantiomer. The enantiomers which are more active in the assays which are described later are preferred forms of the present invention. It will be understood that the present invention includes all mixtures of this active enantiomer with the other enantiomer, for example the racemic mixture.

The active enantiomers may be isolated by separation of racemate for example by fractional crystallization, resolution or HPLC on a chiral column (for example a ChiralpakTM AD 250x50 column). Alternatively the active enantiomers may be made by chiral synthesis from chiral starting materials under conditions which will not cause racemisation or epimerisation, or by derivatisation with a chiral reagent.

The term "prodrug" as used in this specification includes derivatives of the carboxylic acid group which are converted in a mammal, particularly a human, into the carboxylic acid group or a salt or conjugate thereof. It should be understood that, whilst not being bound by theory, it is believed that most of the activity associated with the prodrugs arises from

the activity of the compound of formula I into which the prodrugs are converted. Prodrugs can be prepared by routine methodology well within the capabilities of someone skilled in the art. Various prodrugs of carboxy are known in the art. For examples of such prodrug derivatives, see:

- 5 a) Design of Prodrugs, edited by H. Bundgaard, (Elsevier, 1985) and Methods in Enzymology. 42: 309-396, edited by K. Widder, *et al.* (Academic Press, 1985);
- b) A Textbook of Drug Design and Development, edited by Krogsgaard-Larsen and H. Bundgaard, Chapter 5 "Design and Application of Prodrugs", by H. Bundgaard p.113-191 (1991);
- 10 c) H. Bundgaard, Advanced Drug Delivery Reviews, 8:1-38 (1992);
- d) H. Bundgaard, *et al.*, Journal of Pharmaceutical Sciences, 77:285 (1988); and
- e) N. Kakeya, *et al.*, Chem Pharm Bull, 32:692 (1984).

The above documents a to e are herein incorporated by reference.

In vivo cleavable esters are just one type of prodrug of the parent molecule.

15

The compounds of formula I have activity as medicaments. In particular the compounds of formula I are selective agonists of either PPAR α or PPAR γ , particularly of PPAR α , or are agonists of PPAR α and PPAR γ . The term agonists as used herein, includes partial agonists.

20

Specific compounds of the invention are:

3-{{(3-{{(1,1'-biphenyl-4-ylcarbonyl)amino]methyl}phenyl)amino]methyl}benzoic acid;
2-{{4-(2-oxo-2-{{4-(trifluoromethyl)benzyl}amino}ethyl)phenoxy]methyl}benzoic acid;
25 2-{{(3-{{2-[benzyl(hexyl)amino]-2-oxoethyl}phenoxy)methyl}benzoic acid;
2-{{3-(2-oxo-2-{{4-(trifluoromethyl)benzyl}amino}ethyl)phenoxy]methyl}benzoic acid;
2-{{(4-{{3-[[2-(3,4-dimethoxyphenyl)ethyl](methyl)amino]-3-oxopropyl}phenoxy}-
methyl}benzoic acid;
2-{{(4-{{2-[(4-methyl-2-[4-(trifluoromethyl)phenyl]-1,3-thiazol-5-yl}carbonyl)amino}-
30 ethyl}phenoxy]methyl}benzoic acid;
2-{{4-[2-{{(2,4-difluorophenyl)amino}carbonyl}amino}ethyl]phenoxy]methyl}benzoic
acid;

2-[(4-{2-[(2-methyl-5-phenyl-3-furoyl)amino]ethyl}phenoxy)methyl]benzoic acid;
2-[(4-{2-[(benzylsulfonyl)amino]ethyl}phenoxy)methyl]benzoic acid;
2-[(4-{2-[benzyl(hexyl)amino]-2-oxoethyl}-2-fluorophenoxy)methyl]benzoic acid;
2-[(4-{2-[benzyl(hexyl)amino]-2-oxoethyl}-2-methoxyphenoxy)methyl]benzoic acid;
5 2-[(4-[3-(3,4-dihydroisoquinolin-2(1H)-yl)-3-oxopropyl]phenoxy)methyl]benzoic acid;
2-[(4-{2-[4-(1H-imidazol-1-yl)phenoxy]ethyl}-phenoxy)methyl]benzoic acid;
2-[(4-{2-[4-[(methylsulfonyl)oxy]phenoxy]ethyl}phenoxy)methyl]benzoic acid;
2-[(3-{2-[4-(benzyloxy)phenoxy]ethyl}phenoxy)methyl]benzoic acid;
2-[(3-{2-{4-[(methylsulfonyl)oxy]phenoxy}ethyl}phenoxy)methyl]benzoic acid;
10 2-[(3-[2-(4-hydroxyphenoxy)ethyl]phenoxy)methyl]benzoic acid;
2-[(4-{3-[4-(benzyloxy)phenoxy]propyl}phenoxy)methyl]benzoic acid;
2-[(4-{3-{4-[(methylsulfonyl)oxy]phenoxy}propyl}phenoxy)methyl]benzoic acid;
2-[(4-{3-(4-hydroxyphenoxy)propyl}phenoxy)methyl]benzoic acid;
2-[(4-{3-{[2-(2-ethoxyphenyl)ethyl]amino}-3-oxopropyl}phenoxy)methyl]benzoic acid;
15 2-[(4-{3-[ethyl](2-pyridin-2-ylethyl)amino}-3-oxopropyl}phenoxy)methyl]benzoic acid;
and 2-[(2-{3-{2-[benzyl(hexyl)amino]-2-oxoethoxy}phenyl}ethyl)thio]benzoic acid;
and pharmaceutically acceptable salts, prodrugs, solvates and crystalline forms thereof.

In the present specification the expression "pharmaceutically acceptable salts" is intended to define but is not limited to base salts such as the alkali metal salts, alkaline earth metal salts, ammonium salts, salts with basic amino acids, and salts with organic amines.

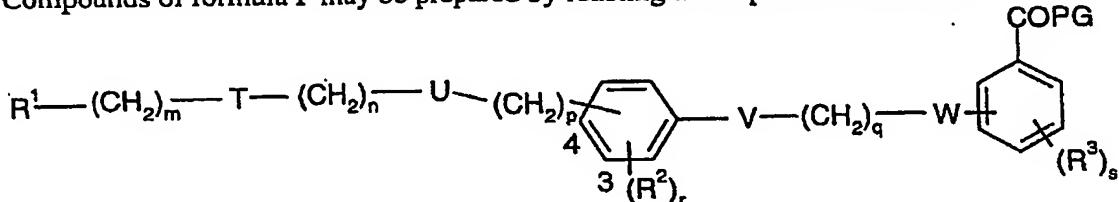
It will also be understood that certain compounds of the present invention may exist in solvated, for example hydrated, as well as unsolvated forms. It is to be understood that the present invention encompasses all such solvated forms. Certain compounds of the present invention may exist as tautomers. It is to be understood that the present invention encompasses all such tautomers.

Methods of preparation

30 The compounds of the invention may be prepared as outlined below. However, the invention is not limited to these methods, the compounds may also be prepared as

described for structurally related compounds in the prior art. The reactions can be carried out according to standard procedures or as described in the experimental section.

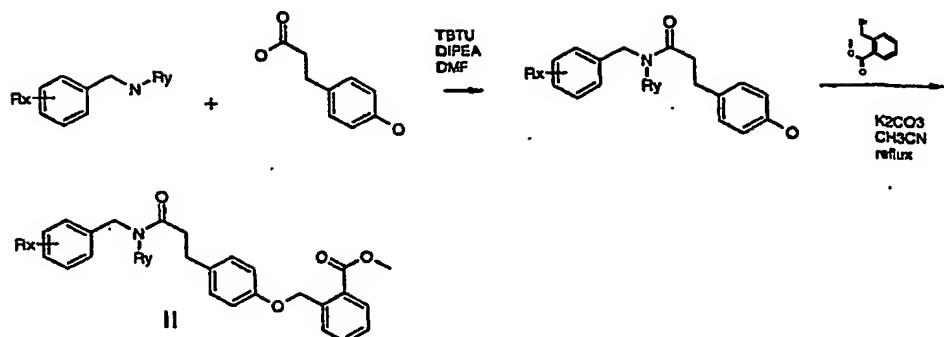
5 Compounds of formula I may be prepared by reacting a compound of formula II



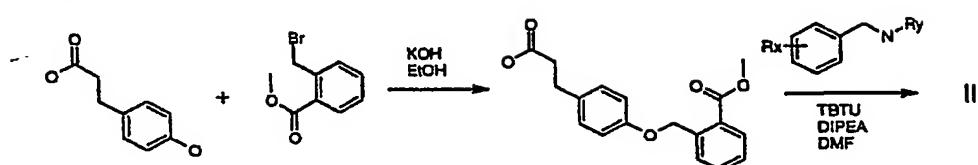
II

in which R^1 , T, U, V, W, R^2 , R^3 , m, n, p, q, r and s are as previously defined and PG represents a protecting group for a carboxylic hydroxy group as described in the standard text "Protective Groups in Organic Synthesis", 2nd Edition (1991) by Greene and Wuts, 10 with a de-protecting agent. The protecting group may also be a resin, such as Wang resin or 2-chlorotriyl chloride resin. Protecting groups may be removed in accordance to techniques which are well known to those skilled in the art. One such protecting group is where PG represents C_{1-6} alkoxy group or an arylalkoxy group eg benzyloxy, such that COPG represents an ester. Such esters can be reacted with a hydrolysing agent, for 15 example lithium hydroxide in the presence of a solvent for example a mixture of THF and water or potassium hydroxide in a C_{1-3} alcohol for example methanol, at a temperature in the range of 0-200°C or by microwave radiation to give compounds of formula I. Compounds of formula II may be prepared according to one the following routes 1 to 5. It will be appreciated by those skilled in the art that methods analogous to those given in 20 routes 1 to 5 may be used to prepare intermediates for compounds of Formula I in which R^1 is a heterocyclic group. Also analogous routes to routes 1 to 5 may be used to prepare compounds of Formula I in which the oxygen atom in the linking chains is replaced by S or NR.

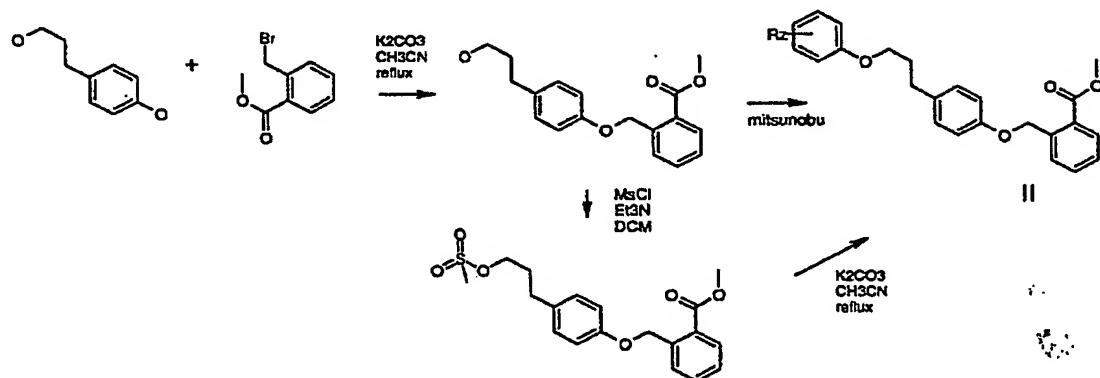
route 1



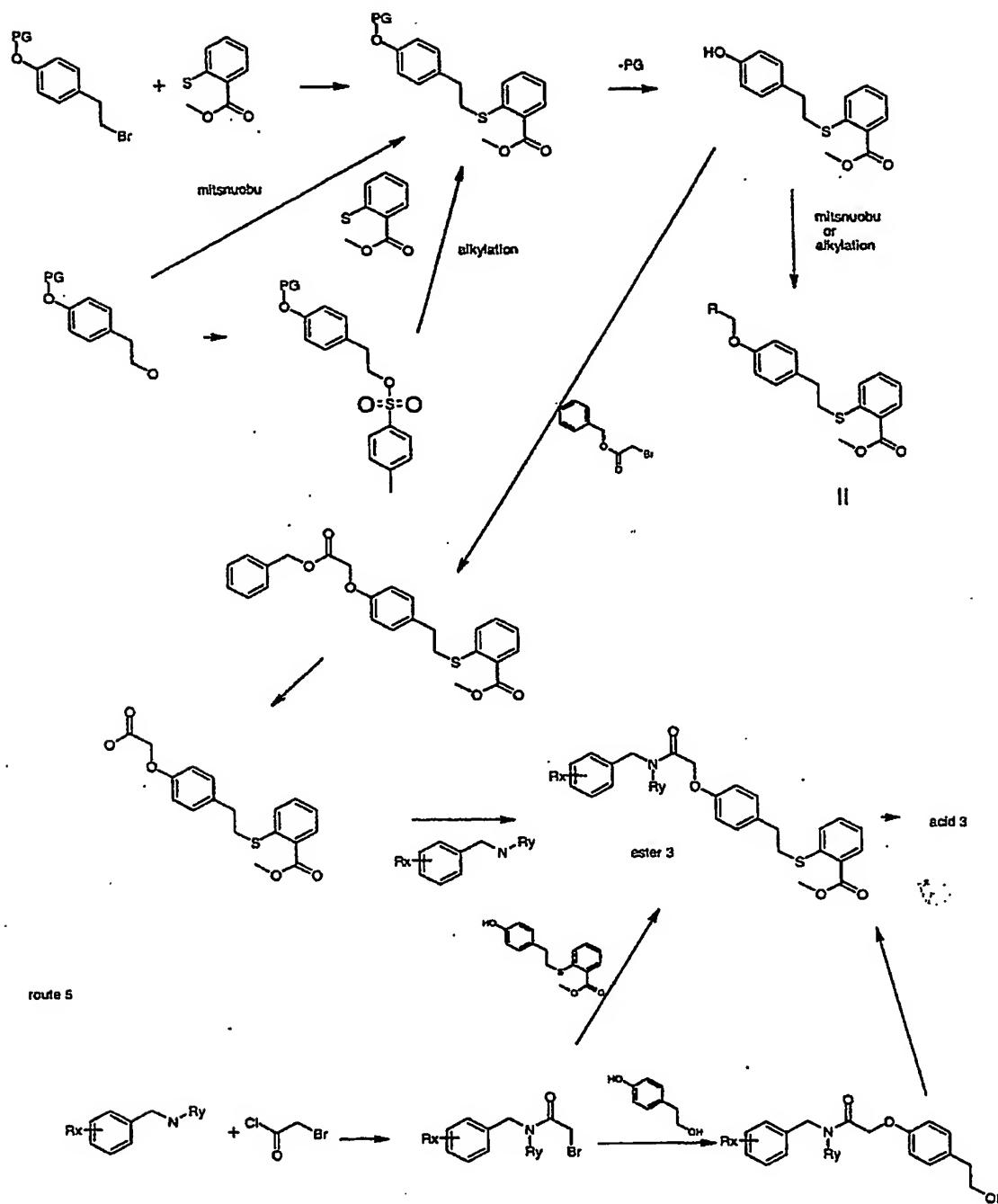
route 2



route 3

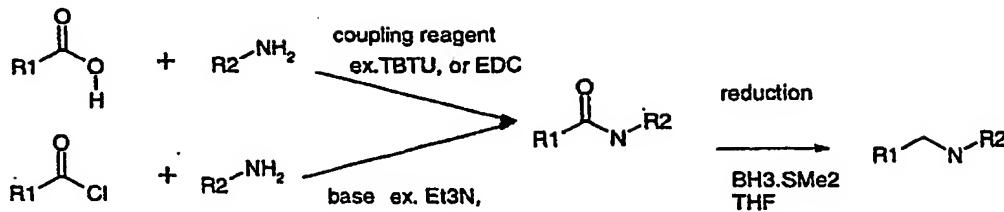


route 4

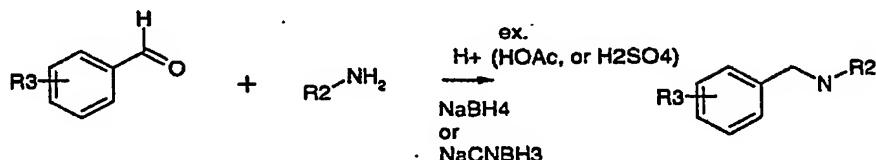


Starting amines may be prepared as described in Ralph N Salvatore et al Tetrahedron, 57, 7785-7811, 2001 or by the methods given below.

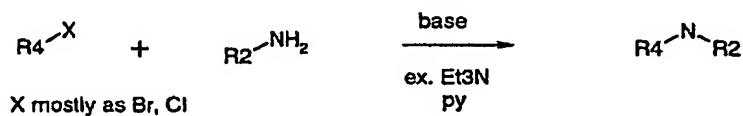
1. Making amide then reduction.



2. reductive amination



3. N-alkylation



Compounds of formula II are believed to be novel and are claimed herein as useful

5 intermediates in the preparation of compounds of formula I.

The compounds of the invention may be isolated from their reaction mixtures using conventional techniques.

10 Persons skilled in the art will appreciate that, in order to obtain compounds of the invention in an alternative and in some occasions, more convenient manner, the individual process steps mentioned hereinbefore may be performed in different order, and/or the individual reactions may be performed at different stage in the overall route (i.e. chemical transformations may be performed upon different intermediates to those associated

15 hereinbefore with a particular reaction).

In any of the preceding methods of preparation, where necessary, hydroxy, amino or other reactive groups may be protected using a protecting group, R^P as described in the standard text "Protective groups in Organic Synthesis", 2nd Edition (1991) by Greene and Wuts. The protecting group may also be a resin, such as Wang resin or 2-chlorotriyl chloride resin. The protection and deprotection of functional groups may take place before or after any of the reaction steps described hereinbefore. Protecting groups may be removed in accordance to techniques which are well known to those skilled in the art.

The expression "inert solvent" refers to a solvent which does not react with the starting materials, reagents, intermediates or products in a manner which adversely affects the yield of the desired product.

Pharmaceutical preparations

The compounds of the invention will normally be administered via the oral, parenteral, intravenous, intramuscular, subcutaneous or in other injectable ways, buccal, rectal, vaginal, transdermal and/or nasal route and/or via inhalation, in the form of pharmaceutical preparations comprising the active ingredient either as a free acid, or a pharmaceutically acceptable organic or inorganic base addition salt, in a pharmaceutically acceptable dosage form. Depending upon the disorder and patient to be treated and the route of administration, the compositions may be administered at varying doses.

Suitable daily doses of the compounds of the invention in therapeutical treatment of humans are about 0.0001-100 mg/kg body weight, preferably 0.001-10 mg/kg body weight.

Oral formulations are preferred particularly tablets or capsules which may be formulated by methods known to those skilled in the art to provide doses of the active compound in the range of 0.5mg to 500mg for example 1 mg, 3 mg, 5 mg, 10 mg, 25mg, 50mg, 100mg and 250mg.

According to a further aspect of the invention there is thus provided a pharmaceutical formulation including any of the compounds of the invention, or pharmaceutically acceptable derivatives thereof, in admixture with pharmaceutically acceptable adjuvants, diluents and/or carriers.

5

Pharmacological properties

The present compounds of formula (I) are useful for the prophylaxis and/or treatment of clinical conditions associated with inherent or induced reduced sensitivity to insulin

10 (insulin resistance) and associated metabolic disorders. These clinical conditions will include, but will not be limited to, general obesity, abdominal obesity, arterial hypertension, hyperinsulinaemia, hyperglycaemia, type 2 diabetes and the dyslipidaemia characteristically appearing with insulin resistance. This dyslipidaemia, also known as the atherogenic lipoprotein profile, phenotype B, is characterised by moderately elevated non-esterified fatty acids, elevated very low density lipoproteins (VLDL) triglyceride rich 15 particles, high Apo B, low high density lipoproteins (HDL) cholesterol, low apoAI particle levels and the presence of small, dense, low density lipoproteins (LDL) particles.

Treatment with the present compounds is expected to lower the cardiovascular morbidity and mortality associated with atherosclerosis due to antidysslipidaemic as well as 20 antiinflammatory properties. The cardiovascular disease conditions include macroangiopathies of various internal organs causing myocardial infarction, congestive heart failure, cerebrovascular disease and peripheral arterial insufficiency of the lower

extremities. Because of their insulin sensitizing effect the compounds of formula I are also expected to prevent or delay the development of type 2 diabetes from the insulin resistance 25 syndrome and diabetes of pregnancy. Therefore the development of long-term complications associated with chronic hyperglycaemia in diabetes mellitus such as the micro-angiopathies causing renal disease, retinal damage and peripheral vascular disease of the lower limbs are expected to be delayed. Furthermore the compounds may be useful in treatment of various conditions outside the cardiovascular system associated with insulin

resistance, like polycystic ovarian syndrome, adipositas, cancer and states of inflammatory disease.

The compounds of the invention may also be combined with other therapeutic agents which are useful in the treatment of disorders associated with the development and progress of atherosclerosis such as hypertension, hyperlipidaemias, dyslipidaemias, diabetes and obesity. In patients with diabetes mellitus the compounds of the invention may also be combined with therapeutic agents used to treat complications related to micro-angiopathies

The compounds of the invention may be used alongside other additional existing therapies for the treatment of type 2 diabetes and its associated complications, these include biguanide drugs, for example metformin, phenformin and buformin, insulin (synthetic insulin analogues, amylin) and oral antihyperglycemics (these are divided into prandial glucose regulators and alpha-glucosidase inhibitors). An example of an alpha-glucosidase inhibitor is acarbose or voglibose or miglitol. An example of a prandial glucose regulator is repaglinide or nateglinide. In addition the combination of the invention may be used in conjunction with another PPAR modulating agent. PPAR modulating agents include but are not limited to thiazolidine-2,4-diones for example troglitazone, ciglitazone, rosiglitazone and pioglitazone. In addition the combination of the invention may be used in conjunction with a sulfonylurea for example: glimepiride, glibenclamide (glyburide), gliclazide, glipizide, gliquidone, chloropropamide, tolbutamide, acetohexamide, glycopyramide, carbutamide, glibenuride, glisoxepid, glybuthiazole, glibuzole, glyhexamide, glymidine, glypinamide, phenbutamide, tolcylamide and tolazamide.

Preferably the sulfonylurea is glimepiride or glibenclamide (glyburide). More preferably the sulfonylurea is glimepiride. Therefore the present invention includes administration of a compound of the present invention in conjunction with one, two or more existing therapies described in this paragraph. The doses of the other existing therapies for the treatment of type 2 diabetes and its associated complications will be those known in the art and approved for use by regulatory bodies for example the FDA and may be found in the Orange Book published by the FDA. Alternatively smaller doses may be used as a result of the benefits derived from the combination.

The present invention also includes a compound of the present invention in combination with a cholesterol-lowering agent. The cholesterol-lowering agents referred to in this application include but are not limited to inhibitors of HMG-CoA reductase (3-hydroxy-3-methylglutaryl coenzyme A reductase). Suitably the HMG-CoA reductase inhibitor is a statin selected from the group consisting of atorvastatin, cerivastatin, fluvastatin, itavastatin, lovastatin, mevastatin, nicostatin, niva-statin, pravastatin and simvastatin, or a pharmaceutically acceptable salt, especially sodium or calcium, or a solvate thereof, or a solvate of such a salt. A particularly preferred statin is, however, a compound with the chemical name (E)-7-[4-(4-fluorophenyl)-6-isopropyl-2-[methyl(methylsulfonyl)-amino]-pyrimidin-5-yl](3R,5S)-3,5-dihydroxyhept-6-enoic acid, [also known as (E)-7-[4-(4-fluorophenyl)-6-isopropyl-2-[N-methyl-N-(methylsulfonyl)-amino]pyrimidin-5-yl](3R,5S)-3,5-dihydroxyhept-6-enoic acid] or a pharmaceutically acceptable salt or solvate thereof, or a solvate of such a salt. The compound (E)-7-[4-(4-fluorophenyl)-6-isopropyl-2-[methyl-(methylsulfonyl)-amino]-pyrimidin-5-yl](3R,5S)-3,5-dihydroxyhept-6-enoic acid, and its calcium and sodium salts are disclosed in European Patent Application, Publication No. EP-A-0521471, and in Bioorganic and Medicinal Chemistry, (1997), 5(2), 437-444. This latter statin is now known under its generic name rosuvastatin.

20 In the present application, the term "cholesterol-lowering agent" also includes chemical modifications of the HMG-CoA reductase inhibitors, such as esters, prodrugs and metabolites, whether active or inactive.

25 The present invention also includes a compound of the present invention in combination with an inhibitor of the ileal bile acid transport system (IBAT inhibitor) for example those described in WO 93/16055, WO 96/16051, WO 94/18183, WO 94/18184, WO 96/05188
WO 96/08484, WO 97/33882, WO 98/07449, WO 98/03818, WO 99/32478, WO
99/64409, WO 00/01687, WO 00/62810, WO 01/66533, WO 02/32428, EP864582,
EP489423, EP549967, EP573848, EP624593, EP624594, EP624595 and EP624596

30 which are hereby incorporated by reference.

The present invention provides a method of treating or preventing insulin resistance (as defined above) comprising the administration of a compound of formula I to a mammal (particularly a human) in need thereof.

5 In a further aspect the present invention provides the use of a compound of formula I in the manufacture of a medicament for the treatment of insulin resistance.

Working examples

10 ^1H NMR and ^{13}C NMR measurements were performed on a Varian Mercury 300 or Varian UNITY plus 400, 500 or 600 spectrometers, operating at ^1H frequencies of 300, 400, 500 and 600 MHz, respectively, and at ^{13}C frequencies of 75, 100, 125 and 150 MHz, respectively. Measurements were made on the delta scale (δ).

15 Unless otherwise stated, chemical shifts are given in ppm with the solvent as internal standard.

Abbreviations

IRS	insulin resistance syndrome
TLC	thin layer chromatography
	HOBtxH ₂ O 1-hydroxybenzotriazole-hydrate
20	DIBAH diisobutylaluminium hydride
	DMSO dimethyl sulfoxide
	EtOAc ethyl acetate
	DMF <i>N,N</i> -dimethylformamide
	THF tetrahydrofuran
25	HPLC high performance liquid chromatography
	MeCN acetonitrile
	TFA trifluoroacetic acid
	Pd/C palladium on charcoal
	HATU O-(7-azabenzotriazolyl-1- yl)-N,N,N',N'-tetramethyluronium
30	hexafluorophosphate
	DCM dichloromethane
	NH ₄ OAc ammonium acetate

MeOH Methanol

DIPEA *N,N*-diisopropylethylamine

DMAP 4-dimethylaminopyridine

Trisamine Tris(hydroxymethyl)aminomethane

5 TBTU O-(benzotriazol-1-yl)-N,N,N',N'-tetramethyluronium tetrafluoroborate

ISOLUTE ® FLASH Si is a silica column suitable for chromatography

Borohydride on polymer support is Borohydride on Amberlite IRA-400 available from Aldrich

EDC 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride

10 NH₄OAc ammonium acetate

LC-MS liquid chromatography- mass spectroscopy

t triplet

s singlet

d doublet

15 q quartet

qvint quintet

m multiplet

br broad

bs broad singlet

20 dm doublet of multiplet

bt broad triplet

dd doublet of doublet

Example 1

25 a) tert-Butyl 3-[(1,1'-biphenyl-4-ylcarbonyl)aminomethyl]phenylcarbamate
Biphenyl-4-carboxylic acid (981 mg, 4.949 mmol) and 3-(aminomethyl)-1-N-boc-aniline (1.0 g, 4.499 mmol) were mixed in DMF (10 ml). Under stirring, benzotriazol-1-yl-oxytrypyrrolidinophosphonium hexafluorophosphate (2.343 g, 4.504 mmol) was added and then N,N-diisopropylethylamine (1.164 g, 9.007 mmol) was added. The mixture was stirred overnight at room temperature. Water and ethyl acetate were added. The organic phase was

washed with water, sodium hydrogencarbonate (sat.) and water (x2) and dried with magnesium sulphate. The solvent was removed. Diethyl ether was added into the residue. The solid product was filtered, washed with little diethyl ether and dried, 1.44g product was obtained. The filtrate was evaporated to dryness. DCM was added to the residue.

5 Filtration gave 0.12 g more solid product. In total 1.56 g desired product was obtained, yield 86%.

¹H NMR (400 MHz, CDCl₃): δ 3.61 (s, 9H), 4.66 (d, 2H), 6.43 (s, br, 1H), 6.50 (s, 1H), 7.06-7.09 (m, 1H), 7.27-7.30 (m, 2H), 7.38-7.50 (m, 4H), 7.61-7.64 (m, 2H), 7.67 (d, 2H) and 7.88 (2H).

10

b) N-(3-Aminobenzyl)-1,1'-biphenyl-4-carboxamide

tert-Butyl 3-[(1,1'-biphenyl-4-ylcarbonyl)amino]methyl phenylcarbamate (250 mg, 0.6 mmol) was dissolved in DCM (10 ml). Trifluoroacetic acid (0.2 ml) was added. The 15 mixture was stirred overnight. HPLC showed that more than 50% of the starting material was not reacted. More trifluoroacetic acid (0.3 ml) was added. The mixture was stirred overnight again. Water was added into the mixture. The phases were not clear. DCM was evaporated in vacuum. Ethyl acetate was added to the residue. The obtained organic phase was washed with water (x3) and dried with magnesium sulphate. The solvent was then 20 evaporated. Crystal product (185 mg) was obtained, yield 99%.

¹H NMR (500 MHz, CD₃OD): δ 4.64 (s, 2H), 7.27 (d, 1H), 7.37-7.40 (m, 2H), 7.45-7.53 (m, 4H), 7.66 (d, 2H), 7.74 (d, 2H) and 7.95 (d, 2H).

c) 3-[(3-[(1,1'-Biphenyl-4-ylcarbonyl)amino]methyl)phenyl]amino]methyl benzoic acid

25

N-(3-aminobenzyl)-1,1'-biphenyl-4-carboxamide (20 mg, 0.07 mmol) was dissolved in acetic acid (0.5 ml). 3-Carboxybenzaldehyde (14 mg, 0.09 mmol) was added and then sodium borohydride (11 mg, 0.28 mmol) was added. The mixture was stirred at room 30 temperature for 2 hours and evaporated to dryness. DCM was added into the residue. The mixture was loaded on a column (ISOLUTE® SI, 500 mg/3 ml). It was eluted with DCM, MeOH/DCM (0.5:99.5) and then MeOH/DCM (1:99). The product fractions were combined and the solvent was removed, Re-chromatography of the residue on a column (

ISOLUTE® SI, 1 g/6 ml) using DCM, MeOH/DCM (0.5:99.5) and then MeOH/DCM (1:99) as eluant gave 9 mg the desired product, yield 31%.

¹H NMR (500 MHz, CD₃OD): δ 4.37 (s, 2H), 4.46 (d, 2H), 6.53 (d, 1H), 6.59-6.61 (m, 2H), 7.04 (t, 1H), 7.30-7.38 (m, 2H), 7.46 (t, 2H), 7.55 (d, 1H), 7.65-7.70 (m, 4H), 7.83 (t, 3H), 8.02 (s, 1H) and 8.80 (br, 1H).

Example 2

10 a) Methyl 2-{{4-(2-oxo-2-{{4-(trifluoromethyl)benzyl}amino}ethyl)phenoxy}methyl}-benzoate

(4-{{2-(Methoxycarbonyl)benzyl}oxy}phenyl)acetic acid (50 mg, 0.167 mmol) was dissolved in DCM (2 ml), 4-(Trifluoromethyl)benzylamine (35 mg, 0.2 mmol) was added, 15 then EDC (38 mg, 0.2 mmol) was added and then DMAP (24.4 mg, 0.2 mmol) was added. The mixture was stirred at room temperature overnight. 1% hydrochloric acid (1 ml) and water (1 ml) was added into the mixture. The two phases were separated using a Whatman Filter Tube. The obtained organic solution was evaporated in vacuum and the solid product (72 mg) was left, yield 95%.

20 ¹H NMR (500 MHz, CDCl₃): δ 3.63 (s, 2H), 3.93 (s, 3H), 4.50 (d, 2H), 5.53 (s, 2H), 5.79 (br, 1H), 7.02 (d, 2H), 7.22 (d, 2H), 7.32 (d, 2H), 7.42 (t, 1H), 7.57-7.61 (m, 3H), 7.77 (d, 1H) and 8.07 (d, 1H).

25 b) 2-{{4-(2-Oxo-2-{{4-(trifluoromethyl)benzyl}amino}ethyl)phenoxy}methyl}benzoic acid

Methyl 2-{{4-(2-oxo-2-{{4-(trifluoromethyl)benzyl}amino}ethyl)phenoxy}methyl}-benzoate (71 mg, 0.155 mmol) in THF (1.5 ml) was cooled in an ice-bath. Lithium hydroxide (7.5 mg, 0.310 mmol) in water (1.5 ml) was dropped in. The cooling-bath was then removed and the mixture was stirred overnight. HPLC showed that the reaction was 30 not complete. More lithium hydroxide (0.2M, 0.5 ml) was added. The reaction mixture was stirred for 4 days more. It was then evaporated in vacuum to remove THF. The residue was acidified with 1% hydrochloric acid, pH=3-4, and then extracted with ethyl acetate. The

organic phase was dried with magnesium sulphate and evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 2g/6 ml) using DCM, MeOH/DCM (0.5:99.5, then 1:99, and then 2:98) as eluant gave 39 mg white solid product, yield 57%.

1 ¹H NMR (500 MHz, CD₃OD): δ 3.50 (s, 2H), 4.42(d, 2H), 5.47 (s, 2H), 6.94 (d, 2H), 7.22 (d, 2H), 7.37-7.40 (m, 3H), 7.53-7.58(m, 3H), 7.69 (d, 1H) and 8.01 (d, 1H).

Example 3

a) (3-{[2-(Methoxycarbonyl)benzyl]oxy}phenyl)acetic acid

10 3-Hydroxyphenylacetic acid (760 mg, (5 mmol) was dissolved in ethanol (99.5%, 20 ml). Potassium hydroxide (560 mg, 10 mmol) was added. The mixture was stirred for 30 minutes. 2-Bromomethylbenzoic acid methyl ester (1.144 g, 5 mmol) was then dropped in. The resulting mixture was heated to reflux for 2 hours and then evaporated in vacuum to dry. Water and ethyl acetate were added into the residue and the phases were separated. The water phase was acidified with 10% hydrochloric acid, pH-5, and then extracted with ethyl acetate. The organic phase was dried (magnesium sulphate) and evaporated in vacuum to dry. Chromatography of the residue on a column (ISOLUTE® SI, 5g/25ml) using DCM, MeOH/DCM (1:99) as eluant gave 213 mg the desired product, yield 14%.

20 ¹H NMR (500 MHz, CDCl₃): δ 3.65 (s, 2H), 3.91 (s, 3H), 5.51 (s, 2H), 6.90-6.96 (m, 3H), 7.27 (t, 1H), 7.39 (t, 1H), 7.57 (t, 1H), 7.77 (d, 1H) and 8.04 (d, 1H).

b) Methyl 2-[(3-{2-[benzyl(hexyl)amino]-2-oxoethyl}phenoxy)methyl]benzoate

25 (3-{[2-(Methoxycarbonyl)benzyl]oxy}phenyl)acetic acid (60 mg, 0.2 mmol) was dissolved in DCM (2 ml), N-hexylbenzylamine (46 mg, 0.24 mmol) was added, then EDC (46 mg, 0.24 mmol) was added and then DMAP (29.3 mg, 0.24 mmol) was added. The mixture was stirred at room temperature overnight. 1% hydrochloric acid (1 ml) and water (1 ml) were added into the mixture. The two phases were separated by using a Whatman Filter Tube. The obtained organic portion was evaporated in vacuum and 59 mg crude oil product was left. It was then used directly in next step.

c) 2-[3-[2-[Benzyl(hexyl)amino]-2-oxoethyl]phenoxy]methylbenzoic acid

5 Methyl 2-[3-[2-[benzyl(hexyl)amino]-2-oxoethyl]phenoxy]methylbenzoate (59 mg, 0.125 mmol) in THF (1 ml) was cooled in an ice-bath. Lithium hydroxide (6 mg, 0.249 mmol) in water (1 ml) was dropped in. The cooling-bath was then removed and the mixture was stirred for 13 days and then evaporated in vacuum to remove THF. The residue was acidified with 1% hydrochloric acid, pH~4, and extracted with ethyl acetate.

10 The organic phase was dried (magnesium sulphate) and evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 1g/6 ml) using DCM and MeOH/DCM (0.5:99.5, then 1:99) as eluant gave 7 mg the desired product, yield 8% (two steps).

15 ¹H NMR (rotamers, 500 MHz, CDCl₃): δ 0.85-0.90 (m, 3H), 1.20-1.30 (m, 6H), 1.45-1.57 (m, 2H), 3.20, 3.40 (t, t, 2H), 3.70, 3.80 (s, s, 2H), 4.51, 4.65 (s, s, 2H), 5.51, 5.52 (s, s, 2H), 6.83-7.00 (m, 3H), 7.14-7.43 (m, 7H), 7.59 (t, 1H), 7.78 (d, 1H) and 8.13 (d, 1H).

Example 4

a) Methyl 2-[3-(2-oxo-2-[4-(trifluoromethyl)benzyl]amino)ethyl]phenoxy]methylbenzoate

20 (3-([2-(Methoxycarbonyl)benzyl]oxy)phenyl)acetic acid (60 mg, 0.2 mmol) was dissolved in DCM (2 ml). 4-(Trifluoromethyl)benzylamine (42 mg, 0.24 mmol) was added, then EDC (46 mg, (0.24 mmol) was added and then DMAP (29.3 mg, 0.24 mmol) was added.

25 The mixture was stirred at room temperature overnight. 1% Hydrochloric acid (1 ml) and water (1 ml) were added to the mixture. The two phases were separated by using a Whatman Filter Tube. The obtained organic portion was evaporated in vacuum and 82 mg solid product was left. It was then used directly in next step.

b) 2-{{3-(2-Oxo-2-{{4-(trifluoromethyl)benzyl}amino}ethyl)phenoxy}methyl}benzoic acid

Methyl 2-{{3-(2-oxo-2-{{4-(trifluoromethyl)benzyl}amino}ethyl)phenoxy}methyl}-benzoate (82 mg, 0.18 mmol) in THF (2 ml) was cooled in an ice-bath. Lithium hydroxide (8.6 mg, 0.36 mmol) in water (1 ml) was dropped in. The cooling-bath was then removed and the mixture was stirred for 7 days and then evaporated in vacuum to remove THF. The residue was acidified with 1% hydrochloric acid, pH~3, and extracted with ethyl acetate. The organic phase was dried (magnesium sulphate) and evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 2g/6 ml) using DCM and MeOH/DCM (1:99, then 2:98) as eluant gave 20 mg the desired product, yield 22.5% (two steps).

¹H NMR (500 MHz, CD₃OD): δ 3.55 (s, 2H), 4.43 (s, 2H), 5.47 (s, 2H), 6.90 (t, 2H), 6.98 (s, 1H), 7.23 (t, 1H), 7.38-7.42 (m, 3H), 7.53-7.59 (m, 3H), 7.70 (d, 1H) and 8.03 (d, 1H).

Example 5

15

a) N-[2-(3,4-dimethoxyphenyl)ethyl]-3-(4-hydroxyphenyl)-N-methylpropanamide

3-(4-Hydroxyphenyl)propionic acid (166.2 mg, 1 mmol) was dissolved in DMF (4 ml). 2-(3,4-Dimethoxyphenyl)-N-methylethylamine (211 mg, 1.05 mmol) was added. The mixture was cooled in an ice-bath. TBTU (337 mg, 1.05 mmol) was added, followed by DIPEA (0.37 ml, 2.1 mmol). The mixture was stirred overnight and the temperature was allowed up to room temperature. Ethyl acetate and sodium hydrogencarbonate aqueous solution (sat.) were added and then the two phases were separated. The water phase was extracted with ethyl acetate. The organic phases were combined and dried (magnesium sulphate) and evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 5g/15 ml) using DCM and then MeOH/DCM (1:99) as eluant gave 333 mg the desired product, yield 97%.

30

¹H NMR (rotamers, 500 MHz, CDCl₃): δ 2.35, 2.59 (t, t, 2H), 2.71-2.80, 2.90 (m, t, 4H), 2.84, 2.97 (s, s, 3H), 3.45, 3.58 (t, t, 2H), 3.84-3.86 (m, 6H), 6.61-6.83 (m, 4H), 6.95 (d, 1H), 7.05 (d, 1H) and 7.50, 7.56 (s, s, 1H).

b) Methyl 2-[(4-{3-[(2-(3,4-dimethoxyphenyl)ethyl](methyl)amino]-3-oxopropyl}phenoxy)methyl]benzoate

N-[2-(3,4-dimethoxyphenyl)ethyl]-3-(4-hydroxyphenyl)-*N*-methylpropanamide (198 mg,

5 0.577 mmol), 2-bromomethyl-benzoic acid methyl ester (139 mg, 0.605 mmol) and potassium carbonate, anhydrous (120 mg, 0.864 mmol) were mixed in acetonitrile (15 ml). The mixture was heated to reflux overnight and then evaporated to dry. Water and ethyl acetate were added and the two phases were separated. The organic phase was dried (magnesium sulphate) and evaporated. Chromatography of the residue on a column 10 (ISOLUTE® SI, 2g/6ml) using heptane/DCM (50:50), then DCM and then MeOH/DCM (0.5:99.5) as eluant gave 172 mg the desired product, yield 61%.

¹H NMR (rotamers, 500 MHz, CDCl₃): 2.34, 2.58 (t, t, 2H), 2.70-2.97 (m, 7H), 3.44, 3.53 (t, t, 2H), 3.84-3.92 (m, 9H), 5.49 (s, br, 2H), 6.61-6.82 (m, 3H), 6.92 (t, 2H), 7.05 (d, 1H), 7.15 (d, 1H), 7.38 (t, 1H), 7.55 (t, 1H), 7.74-7.77 (m, 1H) and 8.03 (d, 1H).

15

c) 2-[(4-{3-[(2-(3,4-Dimethoxyphenyl)ethyl](methyl)amino]-3-oxopropyl}phenoxy)-methyl]benzoic acid

Lithium hydroxide (12 mg, 0.488 mmol) in water (1 ml) was added to methyl 2-[(4-{3-[(2-(3,4-dimethoxyphenyl)ethyl](methyl)amino]-3-oxopropyl}phenoxy)methyl]benzoate (120 mg, 0.244 mmol) dissolved in THF (2 ml). The mixture was then irradiated in a microwave oven (Smith Synthesizer) at 150 °C for 7 minutes and then evaporated to remove THF. The residue was acidified with 1% hydrochloric acid, pH~4, and extracted with ethyl acetate (x2). The organic extracts were combined and washed with brine and dried with

25 magnesium sulphate and then evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 2g/6ml) using DCM, then MeOH/DCM (1:99) as eluant gave 102 mg the desired product, yield 87.5%.

¹H NMR (rotamers, 500 MHz, CDCl₃): δ 2.40, 2.64 (t, t, 2H), 2.73-3.01 (m, 7H), 3.47, 3.63 (t, t, 2H), 3.85-3.88 (m, 6H), 5.56, 5.57 (s, s, 2H), 6.63-6.83 (m, 3H), 6.93-6.97 (m, 2H), 7.07 (d, 1H), 7.17 (d, 1H), 7.41 (t, 1H), 7.57-7.61 (m, 1H), 7.81 (t, 1H) and 8.18 (d, 1H).

Example 6a) Methyl 2-[4-{2-[*tert*-butoxycarbonyl]aminoethyl}phenoxy)methyl]benzoate

5 *tert*-Butyl 2-(4-hydroxyphenyl)ethylcarbamate (3.534 g, 14.9 mmol), 2-bromomethylbenzoic acid methyl ester (3.582 g, 15.6 mmol) and potassium carbonate, anhydrous (3.087 g, 22.3 mmol) were mixed in acetonitrile (50 ml). The mixture was heated to reflux overnight and then evaporated to dry. Water and ethyl acetate were added and the two phases were separated. The organic phase was dried (magnesium sulphate) and evaporated.

10 Chromatography of the residue on a column (ISOLUTE® SI, 20g/70ml) using DCM and then MeOH/DCM (1:99) as eluant gave 5.427 g the desired product, yield 94.5%.

¹H NMR (500 MHz, CDCl₃): δ 1.44 (s, 9H), 2.72 (t, 2H), 3.30-3.35 (m, 2H), 3.86 (s, 3H), 4.87 (s, br, 1H), 5.46 (s, 2H), 6.92 (d, 2H), 7.09 (d, 2H), 7.33 (t, 1H), 7.51 (t, 1H), 7.74 (d, 1H) and 8.00 (d, 1H).

15

b) Methyl 2-[4-(2-aminoethyl)phenoxy)methyl]benzoate hydrochloride

Methyl 2-[4-{2-[*tert*-butoxycarbonyl]aminoethyl}phenoxy)methyl]benzoate (5.1 g, 13.2 mmol) was dissolved in ethyl acetate (50 ml) and it was cooled in an ice-bath.

20 Hydrochloric acid (4M in dioxane, 30 ml, 120 mmol) was added. The cooling bath was removed after 30 minutes. The mixture was stirred for 3 hours more and white precipitates fell out during the time. The reaction mixture was evaporated to dry. Ethyl acetate (20 ml) was added into the residue. It was then filtered. White solid product (3.785 g) was obtained, yield 89%.

25 ¹H NMR (500 MHz, CDCl₃): δ 3.06 (t, 2H), 3.23 (t, 2H), 3.90 (s, 3H), 5.45 (s, 2H), 6.93 (d, 2H), 7.18 (d, 2H), 7.37 (t, 1H), 7.55 (t, 1H), 7.73 (d, 1H), 8.02 (d, 1H) and 8.35 (s, br, 2H).

c) Methyl 2-[4-{2-[{(4-methyl-2-[4-(trifluoromethyl)phenyl]1,3-thiazol-5-yl}carbonyl)aminoethyl}phenoxy]methyl]benzoate

4-Methyl-2-[4-(trifluoromethyl)phenyl]-1,3-thiazole-5-carboxylic acid (50 mg, 0.174 mmol) and methyl 2-[(4-(2-aminoethyl)phenoxy)methyl]benzoate hydrochloride (59 mg, 0.183 mmol) were mixed in DMF (4 ml) and the mixture was then cooled in an ice-bath. TBTU (59 mg, 0.183 mmol) was added and followed by DIPEA (47.2 mg, 0.366 mmol). The mixture was stirred at room temperature overnight. Ethyl acetate and sodium hydrogencarbonate aqueous solution (sat.) were added. The two phases were separated. The organic phase was washed with water and dried (magnesium sulphate) and evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 2g/6ml) using DCM and then MeOH/DCM (0.5:99.5) as eluant gave 59 mg the desired product, white solid, yield 61%.
¹H NMR (400 MHz, CDCl₃): δ 2.62 (s, 3H), 2.87 (t, 2H), 3.67 (dt, 2H), 3.89 (s, 3H), 5.49 (s, 2H), 5.86 (t, 1H), 6.97 (d, 2H), 7.15 (d, 2H), 7.36 (t, 1H), 7.54 (t, 1H), 7.67 (d, 2H), 7.74 (d, 2H) and 7.99-8.03 (m, 3H).

d) 2-[4-{2-[{(4-Methyl-2-[4-(trifluoromethyl)phenyl]1,3-thiazol-5-yl}carbonyl)aminoethyl}phenoxy]methyl]benzoic acid

Lithium hydroxide (4 mg, 0.166 mmol) in water (1 ml) was added into methyl 2-[(4-{2-[(4-methyl-2-[4-(trifluoromethyl)phenyl]1,3-thiazol-5-yl)carbonyl)amino]ethyl}phenoxy)methyl]benzoate (46 mg, 0.083 mmol) dissolved in THF (2 ml). The mixture was then irradiated in a microwave oven (Smith Synthesizer) at 150 °C for 7 minutes and then evaporated to remove THF. The residue was acidified with 1% hydrochloric acid, pH~4, and extracted with ethyl acetate (x2). The organic extracts were combined and dried with magnesium sulphate and then evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 2g/6ml) using DCM, then MeOH/DCM (1:99) as eluant gave 38 mg the desired product, yield 85%.
¹H NMR (400 MHz, THF-d8): δ 2.65 (s, 3H), 2.85 (t, 2H), 3.54 (dt, 2H), 5.52 (s, 2H), 6.94 (d, 2H), 7.17 (d, 2H), 7.36 (t, 1H), 7.41 (t, 1H), 7.53 (t, 1H), 7.75-7.79 (m, 3H), 8.06 (d, 1H) and 8.14 (d, 2H).

Example 7a) Methyl 2-({4-[2-({[(2,4-difluorophenyl)aminolcarbonyl]amino)ethyl]phenoxy}-methyl)benzoate

5

2,4-Difluorophenyl isocyanate (26.5 mg, 0.171 mmol) and methyl 2-{{4-(2-aminoethyl)phenoxy}methyl}benzoate hydrochloride (55 mg, 0.171 mmol) were mixed in DCM (4 ml). PS-DIEA (3.66mmol/g, 140 mg, 0.512 mmol) was added. The mixture was shaken at room temperature overnight. White precipitates were falling out. The mixture was evaporated to dry. The residue (with addition of DCM, a suspension) was loaded on a column (ISOLUTE® SI, 2g/6ml) and eluted with DCM and then MeOH/DCM (1:99). White solid product 51 mg was obtained, yield 68%.

10 ^1H NMR (400 MHz, DMSO-d6): δ 2.65 (t, 2H), 3.26-3.31 (m, 2H), 3.79 (s, 3H), 5.36 (s, 2H), 6.49 (t, 1H), 6.88-6.97 (m, 3H), 7.12-7.22 (m, 3H), 7.44 (t, 1H), 7.58-7.65 (m, 2H), 15 7.88 (d, 1H), 8.01-8.07 (m, 1H) and 8.23 (s, br, 1H).

b) 2-({4-[2-({[(2,4-Difluorophenyl)amino]carbonyl}amino)ethyl]phenoxy}methyl)benzoic acid

20 Lithium hydroxide (3.7 mg, 0.154 mmol) in water (1 ml) was added into methyl 2-{{4-[2-{{[(2,4-difluorophenyl)amino]carbonyl}amino)ethyl]phenoxy}methyl}benzoate (34 mg, 0.077 mmol) dissolved in THF (2 ml). The mixture was then irradiated in a microwave oven (Smith Synthesizer) at 150 °C for 7 minutes and then evaporated to remove THF. The residue was acidified with 1% hydrochloric acid, pH~4, and extracted with ethyl acetate (x2). The organic extracts were combined and dried with magnesium sulphate and then evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 1g/6ml) using MeOH/DCM (1:99, 2:98, 4:96 and then 10: 90) as eluant gave 18 mg the desired product, yield 55%.

25 ^1H NMR (400 MHz, THF-d8): δ 2.76 (t, 2H), 3.39-3.44 (m, 2H), 5.51 (s, 2H), 6.40 (t, 1H), 6.79-6.94 (m, 4H), 7.16 (d, 2H), 7.35 (t, 1H), 7.53 (t, 1H), 7.76 (d, 1H), 7.93 (s, br, 1H), 30 8.06 (d, 1H) and 8.29-8.35 (m, 1H).

Example 8a) Methyl 2-[(4-{2-[2-methyl-5-phenyl-3-furoyl]amino}ethyl)phenoxy)methyl]benzoate

5 2-Methyl-5-phenylfuran-3-carbonyl chloride (36.4 mg, 0.165 mmol) and methyl 2-[(4-(2-aminoethyl)phenoxy)methyl]benzoate hydrochloride (53 mg, 0.165 mmol) were mixed in DCM (4 ml). PS-DIEA (3.66mmol/g, 135 mg, 0.494 mmol) was added. The mixture was shaken at room temperature overnight. LC-MS showed there was only a trace amount of desired product and a big peak of 2-methyl-5-phenyl-3-furoic acid. TBTU (55 mg, 0.17 mmol) was added. The mixture was shaken for 2 hours and filtered. The filtrate was 10 evaporated to dryness. Chromatography of the residue on a column (ISOLUTE® SI, 2g/6ml) using DCM and then MeOH/DCM (0.5:99.5) as eluant gave 34 mg the desired product, yield 44%.

15 ^1H NMR (500 MHz, CDCl_3): δ 2.61 (s, 3H), 2.85 (t, 2H), 3.60-3.66 (m, 2H), 3.89 (s, 3H), 5.49 (s, 2H), 5.82 (t, 1H), 6.55 (s, 1H), 6.96 (d, 2H), 7.15 (d, 2H), 7.25 (t, 1H), 7.36 (t, 3H), 7.52-7.63 (m, 3H), 7.75 (d, 1H) and 8.02 (d, 1H).

b) 2-[(4-{2-[2-Methyl-5-phenyl-3-furoyl]amino}ethyl)phenoxy)methyl]benzoic acid

20 Lithium hydroxide (3.3 mg, 0.136 mmol) in water (1 ml) was added into methyl 2-[(4-(2-[2-methyl-5-phenyl-3-furoyl]amino)ethyl)phenoxy)methyl]benzoate (32 mg, 0.068 mmol) dissolved in THF(2 ml). The mixture was then irradiated in a microwave oven (Smith Synthesizer) at 150 °C for 7 minutes and then evaporated to 25 remove THF. The residue was acidified with 1% hydrochloric acid, pH~4, and extracted with ethyl acetate (x2). The organic extracts were combined and washed with brine and dried with magnesium sulphate and then evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 2g/6ml) using DCM, then MeOH/DCM (1:99 and 2:98) as eluant gave 22 mg the desired product, yield 71%.

30 ^1H NMR (400 MHz, CDCl_3): δ 2.60 (s, 3H), 2.84 (t, 2H), 3.60-3.65 (m, 2H), 5.51 (s, 2H), 5.90 (t, 1H), 6.56 (s, 1H), 6.94 (d, 2H), 7.13 (d, 2H), 7.23 (t, 1H), 7.32-7.40 (m, 3H), 7.55-7.60 (m, 3H), 7.77 (d, 1H) and 8.13 (d, 1H).

Example 9**a) Methyl 2-[(4-(2-[(benzylsulfonyl)amino]ethyl)phenoxy)methyl]benzoate**

5

Alpha-toluenesulfonyl chloride (38 mg, 0.199 mmol) and methyl 2-[(4-(2-aminoethyl)phenoxy)methyl]benzoate hydrochloride (64 mg, 0.199 mmol) were mixed in DCM (3 ml). PS-DIEA (3.66 mmol/g, 272 mg, 0.997 mmol) was added. The mixture was shaken at room temperature over a weekend. It was then loaded on a column (ISOLUTE® SI, 1g/6ml) and eluted with DCM. The product fractions were combined and evaporated.

10 Oil product (17 mg) was obtained, yield 19%.

¹H NMR (500 MHz, CDCl₃): δ 2.74 (t, 2H), 3.19-3.23 (m, 2H), 3.92 (s, 3H), 4.12 (t, 1H), 4.21 (s, 2H), 5.50 (s, 2H), 6.94 (d, 2H), 7.07 (d, 2H), 7.32-7.42 (m, 6H), 7.57 (t, 1H), 7.75 (d, 1H) and 8.05 (d, 1H).

15

b) 2-[(4-(2-[(Benzylsulfonyl)amino]ethyl)phenoxy)methyl]benzoic acid

Lithium hydroxide (2 mg, 0.077 mmol) in water (0.5 ml) was added into methyl 2-[(4-(2-[(benzylsulfonyl)amino]ethyl)phenoxy)methyl]benzoate (17 mg, 0.038 mmol) dissolved in THF (1 ml). The mixture was then irradiated in a microwave oven (Smith Synthesizer) at 150 °C for 7 minutes and then evaporated to remove THF. The residue was acidified with 1% hydrochloric acid, pH~4, and extracted with ethyl acetate (x2). The organic extracts were combined and dried with MgSO₄ and then evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 500 mg/3 ml) using DCM, then MeOH/DCM (0.5:99.5) as eluant gave 10 mg the desired product, yield 61%.

¹H NMR (400 MHz, CDCl₃): δ 2.70 (t, 2H), 3.16-3.21 (m, 2H), 4.18 (s, 2H), 4.29 (t, 1H), 5.48 (s, 2H), 6.91 (d, 2H), 7.05 (d, 2H), 7.29-7.35 (m, 5H), 7.41 (t, 1H), 7.59 (t, 1H), 7.76 (d, 1H) and 8.14 (d, 1H).

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Example 10a) *N*-Benzyl-2-(3-fluoro-4-hydroxyphenyl)-*N*-hexylacetamide

5 3-Fluoro-4-hydroxyphenylacetic acid (170 mg, 0.999 mmol) dissolved in DMF (3 ml) was cooled in an ice-bath. N-Hexylbenzylamine (201 mg, 1.049 mmol) was added and then TBTU (337 mg, 1.049 mmol) followed by DIPEA (407 mg, 3.147 mmol). The mixture was stirred at room temperature overnight and evaporated. Sodium hydrogencarbonate aqueous solution (sat.) was added into the residue. The mixture was then extracted with
10 ethyl acetate (x2). The extracts were combined and washed with water and brine and dried (magnesium sulphate) and evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 5g/15 ml) using DCM and then MeOH/DCM (1:99) as eluant gave 265 mg the desired product, yield 77%.

15 ^1H NMR (rotamers, 400 MHz, CDCl_3): δ 0.83-0.89 (m, 3H), 1.22-1.29 (m, 6H), 1.48-1.58 (m, 2H), 3.21, 3.40 (t, t, 2H), 3.58, 3.68 (s, s, 2H), 4.54, 4.63 (s, s, 2H), 6.72-6.97 (m, 3H), 7.15 (d, 1H), 7.21-7.32 (m, 3H) and 7.35-7.39 (m, 1H).

b) *Methyl* 2-[4-(2-[benzyl(hexyl)amino]-2-oxoethyl)-2-fluorophenoxy]methyl]benzoate

20 *N*-benzyl-2-(3-fluoro-4-hydroxyphenyl)-*N*-hexylacetamide (142 mg, 0.414 mmol), 2-bromomethylbenzoic acid methyl ester (99.4 mg, 0.434 mmol) and potassium carbonate anhydrous (86 mg, 0.620 mmol) were mixed in acetonitrile (5 ml). The mixture was heated to reflux overnight and then evaporated to dry. Ethyl acetate and water were added and the two phases were separated. The organic phase was washed with brine and dried
25 (magnesium sulphate) and evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 2g/6 ml) using DCM, MeOH/DCM (0.5:99.5) as eluant gave 144 mg the desired product, yield 71%.

30 ^1H NMR (rotamers, 400 MHz, CDCl_3): δ 0.83-0.89 (m, 3H), 1.20-1.29 (m, 6H), 1.45-1.58 (m, 2H), 3.18, 3.37 (t, t, 2H), 3.58, 3.69 (s, s, 2H), 3.90 (s, 3H), 4.50, 4.60 (s, s, 2H), 5.53, 5.55 (s, s, 2H), 6.82-7.39 (m, 9H), 7.56 (t, 1H), 7.78-7.82 (m, 1H) and 8.02 (d, 1H).

c) 2-[(4-{2-[Benzyl(hexyl)amino]-2-oxoethyl}-2-fluorophenoxy)methyl]benzoic acid

Methyl 2-[(4-{2-[benzyl(hexyl)amino]-2-oxoethyl}-2-fluorophenoxy)methyl]benzoate (109 mg, 0.222 mmol) was dissolved in THF (2 ml). Lithium hydroxide (10.6 mg, 0.444 mmol) solved in water (1 ml) was added. The mixture was put in a microwave oven (Smith Synthesizer) at 150 °C for 7 minutes. It was then acidified with 1% hydrochloric acid, pH~3, and extracted with ethyl acetate. The organic extract was washed with brine and dried (magnesium sulphate) and evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 2g / 6 ml) using DCM and then MeOH/DCM (0.5: 99.5, then 1:99) as eluant gave 89 mg the desired product, yield 84%.

¹H NMR (rotamers, 300 MHz, CDCl₃): δ 0.84-0.92 (m, 3H), 1.26 (s, br, 6H), 1.48-1.60 (m, 2H), 3.21, 3.41 (t, t, 2H), 3.65, 3.75 (s, s, 2H), 4.55, 4.66 (s, s, 2H), 5.60 (s, 2H), 6.84-7.43 (m, 9H), 7.62 (t, 1H), 7.85 (d, 1H) and 8.16

15 Example 11a) N-Benzyl-N-hexyl-2-(4-hydroxy-3-methoxyphenyl)acetamide

Homovanillic acid (182 mg, 0.999 mmol) dissolved in DMF (3 ml) was cooled in an ice-bath. N-Hexylbenzylamine (201 mg, 1.049 mmol) was added and then TBTU (337 mg, 1.049 mmol) followed by DIPEA (407 mg, 3.147 mmol). The mixture was stirred at room temperature overnight and evaporated. Sodium hydrogencarbonate aqueous solution (sat.) was added into the residue. The mixture was then extracted with ethyl acetate (x2). The extracts were combined and washed with water and brine and dried (magnesium sulphate) and evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 5g/15 ml) using DCM and then MeOH/DCM (1:99) as eluant gave 264 mg the desired product, yield 74%.

¹H NMR (rotamers, 400 MHz, CDCl₃): δ 0.83-0.89 (m, 3H), 1.20-1.29 (m, 6H), 1.44-1.56 (m, 2H), 3.18, 3.37 (t, t, 2H), 3.60, 3.71 (s, s, 2H), 3.81 (s, br, 3H), 4.50, 4.61 (s, s, 2H), 5.98 (s, br, 1H), 6.62-6.85 (m, 3H) and 7.11-7.36 (m, 5H).

b) Methyl 2-[(4-{2-[benzyl(hexyl)amino]-2-oxoethyl}-2-methoxyphenoxy)methyl]benzoate

N-Benzyl-*N*-hexyl-2-(4-hydroxy-3-methoxyphenyl)acetamide (84 mg, 0.236 mmol), 2-bromomethylbenzoic acid methyl ester (57 mg, 0.248 mmol) and potassium carbonate anhydrous (49 mg, 0.355 mmol) were mixed in acetonitrile (5 ml). The mixture was heated to reflux overnight and then evaporated to dry. Ethyl acetate and water were added and the two phases were separated. The organic phase was washed with brine and dried (magnesium sulphate) and evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 2g/6 ml) using DCM, MeOH/DCM (0.5:99.5) as eluant gave 102 mg the desired product yield 86%.

¹H NMR (rotamers, 400 MHz, CDCl₃): δ 0.82-0.88 (m, 3H), 1.18-1.28 (m, 6H), 1.43-1.55 (m, 2H), 3.17, 3.35 (t, t, 2H), 3.61, 3.71 (s, s, 2H), 3.86 (s, 3H), 3.88 (s, 3H), 4.48, 4.60 (s, s, 2H), 5.55, 5.56 (s, s, 2H), 6.61-7.35 (m, 9H), 7.52 (t, 1H), 7.78 (d, 1H) and 8.01 (d, 1H).

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c) 2-[(4-{2-[Benzyl(hexyl)amino]-2-oxoethyl}-2-methoxyphenoxy)methyl]benzoic acid

Methyl 2-[(4-{2-[benzyl(hexyl)amino]-2-oxoethyl}-2-methoxyphenoxy)methyl]benzoate (98 mg, 0.195 mmol) was dissolved in THF (2 ml). Lithium hydroxide (9.3 mg, 0.389 mmol) solved in water (1 ml) was added. The mixture was put in a microwave oven (Smith Synthesizer) at 150 °C for 7 minutes. It was then acidified with 1% hydrochloric acid, pH~3, and extracted with ethyl acetate. The organic extract was washed with brine and dried (magnesium sulphate) and evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 2g / 6 ml) using DCM and then MeOH/DCM (0.5: 99.5, then 1:99) as eluant gave 43 mg the desired product, yield 45%.

¹H NMR (rotamers, 400 MHz, CDCl₃): δ 0.82-0.88 (m, 3H), 1.18-1.28 (m, 6H), 1.44-1.57 (m, 2H), 3.18, 3.37 (t, t, 2H), 3.64, 3.74 (s, s, 2H), 3.86 (s, 3H), 4.50, 4.62 (s, s, 2H), 5.57, 5.58 (s, s, 2H), 6.63-7.39 (m, 9H), 7.56 (t, 1H), 7.80 (d, 1H) and 8.12 (d, 1H).

Example 12a) 4-[3-(3,4-Dihydroisoquinolin-2(1*H*)-yl)-3-oxopropyl]phenol

3-(4-Hydroxyphenyl)propionic acid (202 mg, 1.216 mmol) in DMF (3 ml) was cooled in
5 an ice-bath. 1,2,3,4-Tetrahydroisoquinoline (170 mg, 1.276 mmol) was added and then
TBTU (410 mg, 1.276 mmol) followed by DIPEA (330 mg, 2.553 mmol). The mixture
was stirred at room temperature overnight. Sodium hydrogencarbonate aqueous solution
(sat.) was added. The mixture was extracted with ethyl acetate (x2). The extracts were
combined and dried (magnesium sulphate) and evaporated. Chromatography of the residue
10 on a column (ISOLUTE® SI, 2g/6 ml) using DCM and then MeOH/DCM (1:99) as eluant
gave 303 mg the desired product, yield 89%.

¹H NMR (rotamers, 400 MHz, CDCl₃): δ 2.72-2.77 (m, 2H), 2.83-2.90 (m, 2H), 2.95-3.01
(m, 2H), 3.63, 3.88 (t, t, 2H), 4.57, 4.79 (s, s, 2H), 6.85-6.90 (m, 2H) and 7.07-7.26 (m,
6H).

15

b) Methyl 2-((4-[3-(3,4-dihydroisoquinolin-2(1*H*)-yl)-3-oxopropyl]phenoxy)methyl)-benzoate

4-[3-(3,4-dihydroisoquinolin-2(1*H*)-yl)-3-oxopropyl]phenol (155 mg, 0.551 mmol) was
20 dissolved in acetonitrile (10 ml). 2-Bromomethylbenzoic acid methyl ester (126 mg, 0.551
mmol) was added followed by potassium carbonate anhydrous (114 mg, 0.826 mmol). The
mixture was heated to reflux overnight and then evaporated to dry. Water and ethyl acetate
were added and two phases were separated. The organic phase was dried (magnesium
sulphate) and evaporated. Column chromatography of the residue on silica gel using ethyl
acetate/heptane (40:60) as eluant gave 135 mg the desired product, yield 57%.

¹H NMR (rotamers, 500 MHz, CDCl₃): δ 2.69-2.74 (m, 2H), 2.82-2.87 (m, 2H), 2.95-3.01
(m, 2H), 3.62, 3.85 (t, t, 2H), 3.93 (s, 3H), 4.52, 4.73 (s, s, 2H), 5.48, 5.50 (s, s, 2H), 6.91-
6.95 (m, 2H), 7.03-7.24 (m, 6H), 7.39 (t, 1H), 7.57 (t, 1H), 7.77 (d, 1H) and 8.05 (d, 1H).

c) 2-({4-[3-(3,4-Dihydroisoquinolin-2(1H)-yl)-3-oxopropyl]phenoxy}methyl)benzoic acid

Lithium hydroxide (14.4 mg, 0.6 mmol) dissolved in water (1 ml) was added into 70377 methyl 2-({4-[3-(3,4-dihydroisoquinolin-2(1H)-yl)-3-oxopropyl]phenoxy}methyl)benzoate (129 mg, 0.3 mmol) in THF (2 ml). The mixture was put in a microwave oven (Smith Synthesizer) and irradiated at 150 °C for 7 minutes and then evaporated to remove THF. The residue was acidified with 1% hydrochloric acid, pH~5, and extracted with ethyl acetate (x2). The extracts were combined and dried (magnesium sulphate) and evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 2g/6ml) using DCM, MeOH/DCM (1:99) as eluant gave 111 mg the desired product yield 89%.

¹H NMR (rotamers, 400 MHz, CDCl₃): δ 2.70-2.73 (m, 2H), 2.79-2.83 (m, 2H), 2.92-3.00 (m, 2H), 3.58, 3.84 (t, t, 2H), 4.50, 4.76(s, s, 2H), 5.50, 5.53 (s, s, 2H), 6.87-6.93 (m, 2H), 6.99-7.22 (m, 6H), 7.39 (t, 1H), 7.57 (t, 1H), 7.78 (d, 1H) and 8.16 (d, 1H).

15 Example 13

a) 4-(2-Hydroxyethyl)phenol (2 g, 14.48 mmol) and methyl 2-(bromomethyl)benzoate (3.48 g, 15.20 mmol) were dissolved in acetonitrile (20 ml). Potassium carbonate anhydrous (4.0 g, 28.95 mmol) was added. After stirring at 60° C for three hours PS-trisamine was added (0.2 eq) and the mixture was stirred overnight. The PS-trisamine was filtered off and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic layer was washed with 3 portions of water (3 X 10 ml). The organic phase was dried (MgSO₄) and the solvent was removed by evaporation to give 3.732 g of methyl 2-{{4-(2-hydroxyethyl)phenoxy}methyl}benzoate (yield 90%).

25 ¹HNMR (300 MHz, CDCl₃): δ 2.37 (bs, 1H), 2.8 (t, 2H), 3.8 (bm, 2H), 3.9 (s, 3H), 5.5 (s, 2H), 6.95 (d, 2H), 7.15 (d, 2H), 7.25 (t, 1H), 7.55 (t, 1H), 7.75 (d, 1H), 8.05 (d, 1H)

b) Methyl 2-{{4-(2-hydroxyethyl)phenoxy}methyl}benzoate (1.2 g, 4.19 mmol) was dissolved in dichloromethane (20 ml) and the solution was cooled to -20° C. Triethylamine (0.64 g, 6.29 mmol) was added dropwise, and then methylsulfonyl chloride (0.53 g, 4.61 mmol) was added dropwise. The ice bath was removed and the mixture was stirred at room temperature for one hour. Diethyl ether (5ml) was added and the precipitate was filtered

off. The organic phase was washed with 2 portions of brine, dried ($MgSO_4$) and the solvent was removed by evaporation. The crude was purified by preparative HPLC (starting with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column

5 KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min). The product containing fractions were pooled and the acetonitrile was removed by evaporation. $EtOAc$ (10 ml) was added and the organic phase was washed with two portions of brine and dried ($MgSO_4$). This gave 0.703 g of methyl 2-[(4-{2-[(methylsulfonyl)oxy]ethyl}phenoxy)methyl]benzoate (yield 46%).
10 1H NMR (300 MHz, $CDCl_3$): δ 2.8 (s, 3H), 2.95 (t, 2H), 3.85 (s, 3H), 4.35 (t, 2H), 5.45 (s, 2H), 6.9 (d, 2H), 7.15 (d, 2H), 7.35 (t, 1H), 7.5 (t, 1H), 7.7 (d, 1H), 7.98 (d, 1H)

15 c) Methyl 2-[(4-{2-[(methylsulfonyl)oxy]ethyl}phenoxy)methyl]benzoate (0.2 g, 0.55 mmol) and 4-(1*H*-imidazol-1-yl)phenol (0.11 g, 0.66 mmol) was dissolved in acetonitrile (10 ml) and potassium carbonate (0.09 g, 0.66 mmol) was added. The mixture was stirred over night at 75° C. Remove the acetonitrile by evaporation, dilute with $EtOAc$ (10 ml) and wash the organic phase with Brine three times, dry ($MgSO_4$) and evaporate. This gave 0.268 g of methyl 2-[(4-{2-[4-(1*H*-imidazol-1-yl)phenoxy]ethyl}phenoxy)methyl]benzoate (yield 90%).

20 d) 1H NMR (300 MHz, $CDCl_3$): δ 3.05 (t, 2H), 3.9 (s, 3H), 4.05-4.2 (bm, 2H), 5.55 (s, 2H), 6.9-7.0 (bm, 4H), 7.1-7.4 (bm, 7H), 7.52 (t, 2H), 7.75 (m, 2H), 7.98 (d, 1H)

25 e) Methyl 2-[(4-{2-[4-(1*H*-imidazol-1-yl)phenoxy]ethyl}phenoxy)methyl]benzoate (0.12 g, 0.28 mmol) was dissolved in a mixture of THF/water (7/1, 5 ml) and $LiOH$ (0.03 g, 1.13 mmol) was added. The reaction was performed in a single node microwave oven (7 min, 150° C). Workup by addition of HCl (1 ml, 1M), extract the product by adding two portions of $EtOAc$ (5 ml). The pooled organic phases were dried ($MgSO_4$) and the solvent was removed by evaporation. The crude was purified by preparative HPLC (starting with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min). The product containing fractions were pooled and the acetonitrile was removed by evaporation. $EtOAc$ (10 ml) was added and the

organic phase was washed with two portions of brine and dried ($MgSO_4$). Evaporation gave 6 mg of 2-[(4-{2-[4-(1H-imidazol-1-yl)phenoxy]ethyl}-phenoxy)methyl]benzoic acid (yield 4.7%).

5 1H NMR (500 MHz, $CDCl_3$): δ 3.05 (t, 2H), 4.18 (t, 2H), 5.55 (s, 2H), 6.95 (d, 2H), 7.05 (m, 3H), 7.25 (d, 2H), 7.3-7.45 (bm, 4H), 7.55 (t, 1H), 7.78 (d, 1H), 7.85 (s, 1H), 8.07 (d, 1H).

Example 14

10 a) 4-(2-Hydroxyethyl)phenol (2 g, 14.48 mmol) and methyl 2-(bromomethyl)benzoate (3.48 g, 15.20 mmol) was dissolved in acetonitrile (20 ml). Potassium carbonate anhydrous (4.0 g, 28.95 mmol) was added. After stirring at 60°C for three hours PS-trisamine was added (0.2 eq) and was stirred overnight. The PS-trisamine was filtered off and the 15 acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic layer was washed with 3 portions of water (3 X 10 ml). The organic phase was dried ($MgSO_4$) and the solvent was removed by evaporation to give 3.732 g of methyl 2-[(4-(2-hydroxyethyl)phenoxy)methyl]benzoate (yield 90%).

20 1H NMR (300 MHz, $CDCl_3$): δ 2.37 (bs, 1H), 2.8 (t, 2H), 3.8 (bm, 2H), 3.9 (s, 3H), 5.5 (s, 2H), 6.95 (d, 2H), 7.15 (d, 2H), 7.25 (t, 1H), 7.55 (t, 1H), 7.75 (d, 1H), 8.05 (d, 1H)

25 b) Methyl 2-[(4-(2-hydroxyethyl)phenoxy)methyl]benzoate (1 g, 3.49 mmol), 4-(benzyloxy)phenol (0.7 g, 3.49 mmol) and triphenylphosphine (1.01 g, 3.84 mmol) was added to a dry round bottomed flask fitted with a septum. N_2 was flushed through the flask for 5 minutes followed by the addition of dry toluene (30 ml) and diisopropylazodicarboxylate (0.78 g, 3.84 mmol). The reaction mixture was stirred at 55°C overnight. The solvent was removed by evaporation and the crude material was purified by 30 preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40

ml/min). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried ($MgSO_4$). After removing the solvent by evaporation, 0.7g of methyl 2-[(4-{2-[4-(benzyloxy)phenoxy]ethyl}phenoxy)methyl]benzoate was isolated (yield 42.8%).

¹HNMR (300 MHz, $CDCl_3$): δ 3.05 (t, 2H), 3.95 (s, 3H), 4.07 (t, 2H), 5.03 (s, 2H), 5.55 (s, 2H), 6.85 (d, 2H), 6.95 (d, 2H), 6.98 (d, 2H), 7.23 (d, 2H), 7.3-7.5 (bm, 6H), 7.6 (t, 1H), 7.8 (d, 1H), 8.07 (d, 1H).

c) Methyl 2-[(4-{2-[4-(benzyloxy)phenoxy]ethyl}phenoxy)methyl]benzoate (0.80 g 1.71 mmol), borontrifluoride etherate (2.42 g, 17.09 mmol) and dimethyl sulfide (1.27 g, 20.51 mmol) were dissolved in dichloromethane (25 ml). The mixture was stirred for 6 hours at room temperature. EtOAc (20 ml) was added and the mixture was washed with three portions of water, the organic layer dried ($MgSO_4$) and the solvent removed by evaporation.

¹HNMR (300 MHz, $CDCl_3$): δ 3.0 (t, 2H), 3.9 (s, 3H), 4.1 (t, 2), 5.5 (s, 2H), 5.55 (s, 2H), 6.8 (bm, 4H), 7.0 (d, 2H), 7.2 (d, 2H), 7.4 (t, 1H), 7.55 (t, 1H), 7.8 (d, 1H), 8.05 (d,

d) Methyl 2-[(4-[2-(4-hydroxyphenoxy)ethyl]phenoxy)methyl]benzoate (0.547 g, 1.45 mmol) was dissolved in dichloromethane (10 ml) and the solution was cooled to -20°C. Triethylamine (0.22 g, 2.17 mmol) was added dropwise followed by the dropwise addition of methylsulfonyl chloride (0.18 g, 1.59 mmol). The ice bath was removed and the mixture was stirred overnight at room temperature. Excess of triethylamine was removed by addition of diethyl ether (5 ml) and filtering off the precipitate. The organic phase was washed with three portions of brine (10 ml) and dried ($MgSO_4$). The solvent was removed by evaporation, the crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the

organic phase was washed with two portions of brine and dried ($MgSO_4$). Removing the solvent by evaporation gave 0.317g of methyl 2-([4-(2-{4-[(methylsulfonyl)oxy]phenoxy}ethyl)phenoxy]methyl)benzoate (yield 48%).

5 1H NMR (500 MHz, $CDCl_3$): δ 3.07 (t, 2H), 3.15 (s, 3H), 3.95 (s, 3H), 4.17 (t, 2H), 5.55 (s, 2H), 6.95 (d, 2H), 7.0 (d, 2H), 7.23 (m, 4H), 7.42 (t, 1H), 7.6 (t, 1H), 7.8 (d, 1H), 8.08 (d, 1H).

10 e) Methyl 2-([4-(2-{4-[(methylsulfonyl)oxy]phenoxy}ethyl)phenoxy]methyl)benzoate (0.18 g, 0.38 mmol) was dissolved in a mixture of tetrahydrofuran and water 7:1 (5 ml). The reaction was performed in a single node microwave oven (150° C in 7 minutes). The mixture was diluted with HCl (2 ml, 1 M) and the organic phase was isolated. The crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of 15 acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried ($MgSO_4$). Removing the solvent by evaporation gave 13 mg of 2-([4-(2-{4-[(methylsulfonyl)oxy]phenoxy}ethyl)phenoxy]- 20 methyl)benzoic acid (yield 7.7%).

15 1H NMR (300 MHz, $CDCl_3$): δ 3.07 (t, 2H), 3.15 (s, 3H), 4.17 (t, 2H) 5.6 (s, 2H), 6.95 (d, 2H), 7.0 (d, 2H), 7.23 (m, 4H), 7.42 (t, 1H), 7.65 (t, 1H), 7.82 (d, 1H), 8.2 (d, 1H).

25 Example 15

a) 3-(2-Hydroxyethyl)phenol (1.0 g, 7.24 mmol) and methyl 2-(bromomethyl)benzoate (1.74 g, 7.6 mmol) were dissolved in acetonitrile (10 ml). Potassium carbonate anhydrous (2.0 g, 14.48 mmol) was added. After stirring at 60°C for three hours PS-trisamine was added (0.3 eq) and was stirred overnight. The PS-trisamine was filtered off and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic layer was washed with 3 portions of water (3 X 10 ml). The organic phase was dried ($MgSO_4$)

and the solvent was removed by evaporation to give 1.99 g of methyl 2-[(3-(2-hydroxyethyl)phenoxy)methyl]benzoate (yield 90%).

¹HNMR (300 MHz, CDCl₃): δ 2.95 (t, 2H), 3.45 (s, 1H), 3.9 – 4.0 (bm, 5H), 5.58 (s, 2H),

5 6.95 (m, 2H), 7.05 (s, 1H), 7.3 (t, 1H), 7.45 (t, 1H), 7.6 (t, 1H), 7.85 (d, 1H), 8.05 (d, 1H)

b) Methyl 2-[(3-(2-hydroxyethyl)phenoxy)methyl]benzoate (0.5g, 1.75 mmol), 4-(benzyloxy)phenol (0.35 g, 1.75 mmol) and triphenylphosphine (0.5 g, 1.92 mmol) were added to a dry roundbottomed flask and fitted with a septum. Dry toluene (10 ml) was added and N₂ was flushed through the mixture for 5 minutes. Diisopropyl (E)-diazene-1,2-dicarboxylate (0.39 g, 1.92 mmol) was added dropwise and the solution was stirred at room temperature. After three hour, another equivalent of reagents was added and stirred for one hour. After removing the solvent by evaporation the crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min). The product containing fractions was pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried (MgSO₄). Removing the solvent by evaporation gave 0.319 g

10 methyl 2-[(3-(2-[4-(benzyloxy)phenoxy]ethyl)phenoxy)methyl]benzoate (yield 39%).

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¹HNMR (300 MHz, CDCl₃): δ 3.15 (t, 2H), 3.95 (s, 3H), 4.2 (t, 2H), 5.07 (s, 2H), 5.6 (s, 2H), 6.9 – 7.1 (bm, 7H), 7.3 – 7.55 (bm, 7H), 7.6 (t, 1H), 7.85 (d, 1H), 8.05 (d, 1H)

c) Methyl 2-[(3-(2-[4-(benzyloxy)phenoxy]ethyl)phenoxy)methyl]benzoate (20 mg, 0.043 mmol) was dissolved in a mixture of THF/H₂O (7/1, 3 ml) and LiOH (4.1 mg, 0.17 mmol) was added. The reaction was performed in a single node microwave oven (7 min, 150° C). The mixture was acidified (HCl, 1 ml, 1M) and the water phase washed with two portions of EtOAc (3 X 5 ml). After removing the solvent by evaporation the crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40

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ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried ($MgSO_4$). Removing the solvent by evaporation gave 19 mg of 2-[(3-{2-[4-(benzyloxy)phenoxy]ethyl}phenoxy)methyl]benzoic acid (yield 98 %).

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1H NMR (300 MHz, $CDCl_3$): δ 3.1 (t, 2H), 4.15 (t, 2H), 5.03 (s, 2H), 5.57 (s, 2H), 6.8 – 7.0 (bm, 7H), 7.2 – 7.5 (bm, 7H), 7.6 (t, 1H), 7.85 (d, 1H), 8.15 (d, 1H)

10 Example 16

a) 3-(2-Hydroxyethyl)phenol (1.0 g, 7.24 mmol) and methyl 2-(bromomethyl)benzoate (1.74 g, 7.6 mmol) were dissolved in acetonitrile (10 ml). Potassium carbonate anhydrous (2.0 g, 14.48 mmol) was added. After stirring at 60°C for three hours PS-trisamine was 15 added (0.3 eq) and was stirred overnight. The PS-trisamine was filtered off and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic layer was washed with 3 portions of water (3 X 10 ml). The organic phase was dried ($MgSO_4$) and the solvent was removed by evaporation to give 1.99 g of methyl 2-[(3-(2-hydroxyethyl)phenoxy)methyl]benzoate (yield 90%).

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1H NMR (300 MHz, $CDCl_3$): δ 2.95 (t, 2H), 3.45 (s, 1H), 3.9 – 4.0 (bm, 5H), 5.58 (s, 2H), 6.95 (m, 2H), 7.05 (s, 1H), 7.3 (t, 1H), 7.45 (t, 1H), 7.6 (t, 1H), 7.85 (d, 1H), 8.05 (d, 1H)

b) Methyl 2-[(3-(2-hydroxyethyl)phenoxy)methyl]benzoate (0.5g, 1.75 mmol), 4-(benzyloxy)phenol (0.35 g, 1.75 mmol) and triphenylphosphine (0.5 g, 1.92 mmol) was 25 added to a dry round bottomed flask and fitted with septum. Dry toluene (10 ml) was added and N_2 was flushed through the mixture for 5 minutes. Diisopropyl (E)-diazene-1,2-dicarboxylate (0.39 g, 1.92 mmol) was added dropwise and the solution was stirred at room temperature. After three hours another equivalent of reagents was added and stirred 30 for one hour. After removing the solvent by evaporation the crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90

and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried ($MgSO_4$). Removing the solvent by evaporation gave 0.319 g
5 methyl 2-[(3-{2-[4-(benzyloxy)phenoxy]ethyl}phenoxy)methyl]benzoate (yield 39%).

¹HNMR (300 MHz, $CDCl_3$): δ 3.15 (t, 2H), 3.95 (s, 3H), 4.2 (t, 2H), 5.07 (s, 2H), 5.6 (s, 2H), 6.9 – 7.1 (bm, 7H), 7.3 – 7.55 (bm, 7H), 7.6 (t, 1H), 7.85 (d, 1H), 8.05 (d, 1H)

10 c) Methyl 2-[(3-{2-[4-(benzyloxy)phenoxy]ethyl}phenoxy)methyl]benzoate (0.275 g, 0.59 mmol) was dissolved in dichloromethane (10 ml), dimethylsulfide (0.44 g, 7.0 mmol) and borontrifloureide eterate (0.83 g, 5.87 mmol) was added and the mixture was stirred at room temperature for six hours. EtOAc (10 ml) was added and the organic phase was washed with water (3 X 10 ml), dried ($MgSO_4$) and the solvent was removed by
15 evaporation and the crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions was pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the
20 organic phase was washed with two portions of brine and dried ($MgSO_4$). After removing the solvent by evaporation 0.096 gram of methyl 2-[(3-[2-(4-hydroxyphenoxy)ethyl]-phenoxy}methyl]benzoate was obtained (yield 43.2%). This product was used directly in the next step.

25 d) Methyl 2-[(3-[2-(4-hydroxyphenoxy)ethyl]phenoxy)methyl]benzoate (0.096 g, 0.25 mmol) was dissolved in dichloromethane (10 ml) and cooled to -20° C. Triethylamine (0.039g, 0.38 mmol) was added drop wise and methanesulfonyl chloride (0.032 g, 0.28 mmol) was added drop wise. The ice bath was removed and the mixture was wormed to room temperature. Diethyl ether (5 ml) was added and the precipitate filtered off. The
30 organic phase was washed with two portions of brine (5 ml) and dried ($MgSO_4$). Removing The solvent was removed by evaporation giving 0.109 gram of methyl 2-[(3-(2-{4-[(methylsulfonyl)oxy]phenoxy}-ethyl)phenoxy)methyl]benzoate (yield 94.1%).

¹HNMR (500 MHz, CDCl₃): δ 3.15 (m, 5H), 3.95 (s, 3H), 4.2 (t, 2H), 5.55 (s, 2H), 6.95 (s, 4H), 7.0 (s, 1H), 7.25 (d, 2H), 7.3 (t, 1H), 7.42 (t, 1H), 7.6 (t, 1H), 7.8 (d, 1H), 8.1 (d, 1H).

5 e) Methyl 2-{[3-(2-{4-[(methylsulfonyl)oxy]phenoxy}ethyl)phenoxy]methyl}benzoate (0.109 g, 0.24 mmol) was dissolved in a mixture of THF/water (7/1, 2.5 ml). Lithium hydroxide (23 mg, 0.96 mmol) was added. The reaction was performed in a single node microwave oven (7 min, 150° C). The mixture was acidified (HCl, 1 ml, 1 M) and the water phase was extracted with two portions of EtOAc (2 X 5 ml). The organic phases 10 were combined, dried (MgSO₄) and the solvent was removed by evaporation and gave 17 mg of 2-{[3-(2-{4-[(methylsulfonyl)oxy]phenoxy}ethyl)phenoxy]methyl}benzoic acid (yield 16%).

¹HNMR (500 MHz, CDCl₃): δ 3.15 (m, 5H), 4.2 (t, 2H), 5.55 (s, 2H), 6.95 (s, 4H), 7.0 (s, 1H), 7.25 (d, 2H), 7.3 (t, 1H), 7.42 (t, 1H), 7.6 (t, 1H), 7.8 (d, 1H), 8.1 (d, 1H).

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Example 17

a) 3-(2-Hydroxyethyl)phenol (1.0 g, 7.24 mmol) and methyl 2-(bromomethyl)benzoate (1.74 g, 7.6 mmol) was dissolved in acetonitrile (10 ml). Potassium carbonate anhydrous 20 (2.0 g, 14.48 mmol) was added. After stirring at 60°C for three hours PS-trisamine was added (0.3 eq) and was stirred overnight. The PS-trisamine was filtered off and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic layer was washed with 3 portions of water (3 X 10 ml). The organic phase was dried (MgSO₄) and the solvent was removed by evaporation to give 1.99 g of methyl 2-{[3-(2- 25 hydroxyethyl)phenoxy]methyl}benzoate (yield 90%).

¹HNMR (300 MHz, CDCl₃): δ 2.95 (t, 2H), 3.45 (s, 1H), 3.9 – 4.0 (bm, 5H), 5.58 (s, 2H), 6.95 (m, 2H), 7.05 (s, 1H), 7.3 (t, 1H), 7.45 (t, 1H), 7.6 (t, 1H), 7.85 (d, 1H), 8.05 (d, 1H)

30 b) Methyl 2-{[3-(2-hydroxyethyl)phenoxy]methyl}benzoate (0.5g, 1.75 mmol), 4-(benzyloxy)phenol (0.35 g, 1.75 mmol) and triphenylphosphine (0.5 g, 1.92 mmol) was added to a dry round bottomed flask and fitted with septum. Dry toluene (10 ml) was

added and N_2 was flushed through the mixture for 5 minutes. Diisopropyl (*E*)-diazene-1,2-dicarboxylate (0.39 g, 1.92 mmol) was added dropwise and the solution was stirred at room temperature. After three hours another equivalent of reagents was added and stirred for one hour. After removing the solvent by evaporation the crude was purified by 5 preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two 10 portions of brine and dried ($MgSO_4$). Removing the solvent by evaporation gave 0.319 g methyl 2-[(3-{2-[4-(benzyloxy)phenoxy]ethyl}phenoxy)methyl]benzoate (yield 39%).

15 1H NMR (300 MHz, $CDCl_3$): δ 3.15 (t, 2H), 3.95 (s, 3H), 4.2 (t, 2H), 5.07 (s, 2H), 5.6 (s, 2H), 6.9 – 7.1 (bm, 7H), 7.3 – 7.55 (bm, 7H), 7.6 (t, 1H), 7.85 (d, 1H), 8.05 (d, 1H)

15 c) Methyl 2-[(3-{2-[4-(benzyloxy)phenoxy]ethyl}phenoxy)methyl]benzoate (0.275 g, 0.59 mmol) was dissolved in dichloromethane (10 ml), dimethyl sulfide (0.44 g, 7.0 mmol) and borontrifluoride etherate (0.83 g, 5.87 mmol) was added and the mixture was stirred at room temperature for six hours. EtOAc (10 ml) was added and the organic phase was 20 washed with water (3 X 10 ml), dried ($MgSO_4$) and the solvent was removed by evaporation and the crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were 25 pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried ($MgSO_4$). After removing the solvent by evaporation 0.096 gram of methyl 2-[(3-[2-(4-hydroxyphenoxy)ethyl]phenoxy)methyl]benzoate was obtained (yield 43.2%). This product was used directly in the next step.

30 d) Methyl 2-[(3-[2-(4-hydroxyphenoxy)ethyl]phenoxy)methyl]benzoate (0.096 g, 0.25 mmol) was dissolved in dichloromethane (10 ml) and cooled to -20° C. Triethylamine

(0.039g, 0.38 mmol) was added drop wise and methanesulfonyl chloride (0.032 g, 0.28 mmol) was added drop wise. The ice bath was removed and the mixture was wormed to room temperature. Add diethyleter (5 ml) and filter of the precipitate, wash the organic phase with two potions of brine (5 ml) and dry (MgSO_4). Removing the solvent by evaporation gave 0.109 gram of methyl 2-[(3-(2-(4-[(methylsulfonyl)oxy]phenoxy)ethyl)-phenoxy)methyl]benzoate (yield 94.1%).

¹⁰ ^1H NMR (500 MHz, CDCl_3): δ 3.15 (m, 5H), 3.95 (s, 3H), 4.2 (t, 2H), 5.55 (s, 2H), 6.95 (s, 4H), 7.0 (s, 1H), 7.25 (d, 2H), 7.3 (t, 1H), 7.42 (t, 1H), 7.6 (t, 1H), 7.8 (d, 1H), 8.1 (d, 1H).

¹⁵ f) Methyl 2-[(3-(2-(4-[(methylsulfonyl)oxy]phenoxy)ethyl)phenoxy)methyl]benzoate (0.109 g, 0.24 mmol) was dissolved in a mixture of THF/water (7/1, 2.5 ml). Lithium hydroxide (23 mg, 0.96 mmol) was added. The reaction was performed in a single node microwave oven (7 min, 150° C). The mixture was acidified (HCl , 1 ml, 1 M) and the water phase was extracted with two potions of EtOAc. The organic phases were combined, dried (MgSO_4) and the solvent was removed by evaporation and gave 16 mg of 2-[(3-[2-(4-hydroxyphenoxy)ethyl]phenoxy)methyl]benzoic acid (yield 18%)

²⁰ ^1H NMR (400 MHz, CDCl_3): δ 3.0 (t, 2H), 4.1 (t, 2H), 5.55 (s, 2H), 6.7 (m, 4H), 6.8-6.95 (bm, 3H), 7.4 (m, 1H), 7.6 (m, 1H), 7.8 (m, 1H), 8.15 (m, 1H).

Example 18

²⁵ a) 4-(3-Hydroxypropyl)phenol (1.0 g, 6.57 mmol) and methyl 2-(bromomethyl)benzoate (1.66 g, 7.23 mmol) was dissolved in acetonitrile (10 ml). Potassium carbonate (1.82 g, 13.14 mmol) was added and the mixture was stirred at 60° C for three hours. Polymersupported trisamine (0.3 eqv) was added and the solution was stirred at room temperature over night. The PS-trisamine was filtered of and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic layer was washed with 3 potions of water (3 X 10 ml). The organic phase was dried (MgSO_4) and the solvent was removed by evaporation to give 1.66 gram of methyl 2-[(4-(3-hydroxypropyl)phenoxy)-methyl]benzoate (yield 84.2%).

¹HNMR (500 MHz, CDCl₃): δ 1.9 (m, 2H), 2.65 (t, 2H), 3.25 (s, 1H), 3.65 (t, 2H), 3.85 (s, 3H), 5.45 (s, 2H), 6.95 (d, 2H), 7.15 (d, 2H), 7.35 (t, 1H), 7.5 (t, 1H), 7.75 (d, 1H), 8.05 (d, 1H).

5 b) Methyl 2-[(4-(3-hydroxypropyl)phenoxy)methyl]benzoate (0.50 g, 1.66 mmol) and 4-(benzyloxy)phenol (0.33 g, 1.66 mmol) was added to a dry round bottomed flask and fitted with septa. Dry toluene (10 ml) was added and N₂ was flushed through the mixture for 5 minutes. (Tributylphosphoranylidene)acetonitrile (0.80 g, 3.33 mmol) was added dropwise and the reaction was performed in a single node microwave oven. After removing the solvent by evaporation the crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried (MgSO₄). Removing the solvent by evaporation gave 0.515 gram of methyl 2-[(4-(3-[4-(benzyloxy)phenoxy]-propyl)phenoxy)methyl]benzoate (yield 64.1%).

20 ¹HNMR (500 MHz, CDCl₃): δ 2.15 (m, 2H), 2.85 (t, 2H), 4.0 (m, 5H), 5.1 (s, 2H), 5.6 (s, 2H), 6.9-7.1 (bm, 6H), 7.22 (d, 2H), 7.35-7.55 (bm, 6H), 7.62 (t, 1H), 7.9 (d, 1H), 8.15 (d, 1H).

25 c) Methyl 2-[(4-(3-[4-(benzyloxy)phenoxy]propyl)phenoxy)methyl]benzoate (0.047 g, 0.097 mmol) was dissolved in a mixture of THF/water (7/1, 2 ml) and lithium hydroxide (9.3 mg, 0.39 mmol) was added. The reaction was performed in a single node microwave oven (7 min, 150° C). The reaction mixture was acidified (HCl, 1 M, 1 ml) and the water phase was washed with two portions of EtOAc (2 X 5 ml). The organic phases were combined, dried (MgSO₄) and the solvent was removed by evaporation. The crude product was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X

250 mm, flow 40 ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried ($MgSO_4$). Removing the solvent by evaporation gave 2 mg of 2-[(4-{3-[4-(benzyloxy)phenoxy]propyl}phenoxy)methyl]-
5 benzoic acid (yield 4.4%).

¹HNMR (500 MHz, $CDCl_3$): δ 2.05 (m, 2H), 2.75 (t, 2H), 4.0 (t, 2H), 5.05 (s, 2H), 5.6 (s, 2H), 6.8-7.0 (bm, 6H), 7.15 (d, 2H), 7.35-7.55 (bm, 6H), 7.55 (t, 1H), 7.8 (d, 1H), 8.15 (d, 1H).

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Example 19

a) 4-(3-Hydroxypropyl)phenol (1.0 g, 6.57 mmol) and methyl 2-(bromomethyl)benzoate (1.66 g, 7.23 mmol) was dissolved in acetonitrile (10 ml). Potassium carbonate (1.82 g, 15 13.14 mmol) was added and the mixture was stirred at 60° C for three hours. Polymersupported trisamine (0.3 eqv) was added and the solution was stirred at room temperature overnight. The PS-trisamine was filtered off and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic layer was washed with 3 portions of water (3 X 10 ml). The organic phase was dried ($MgSO_4$) and the solvent was 20 removed by evaporation to give 1.66 gram of methyl 2-{{4-(3-hydroxypropyl)phenoxy}-methyl}benzoate (yield 84.2%).

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¹HNMR (500 MHz, $CDCl_3$): δ 1.9 (m, 2H), 2.65 (t, 2H), 3.25 (s, 1H), 3.65 (t, 2H), 3.85 (s, 3H), 5.45 (s, 2H), 6.95 (d, 2H), 7.15 (d, 2H), 7.35 (t, 1H), 7.5 (t, 1H), 7.75 (d, 1H), 8.05 (d, 1H).

30

b) Methyl 2-{{4-(3-hydroxypropyl)phenoxy}methyl}benzoate (0.50 g, 1.66 mmol) and 4-(benzyloxy)phenol (0.33 g, 1.66 mmol) was added to a dry round bottomed flask and fitted with septa. Dry toluene (10 ml) was added and N_2 was flushed through the mixture for 5 minutes. (Tributylphosphoranylidene)acetonitrile (0.80 g, 3.33 mmol) was added dropwise and the reaction was performed in a single node microwave oven.. After removing the solvent by evaporation the crude was purified by preparative HPLC (started with

acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried (MgSO_4). Removing the solvent by evaporation gave 0.515 gram of methyl 2-[(4-{3-[4-(benzyloxy)phenoxy]-propyl}phenoxy)methyl]benzoate (yield 64.1%).

10 ^1H NMR (500 MHz, CDCl_3): δ 2.15 (m, 2H), 2.85 (t, 2H), 4.0 (m, 5H), 5.1 (s, 2H), 5.6 (s, 2H), 6.9-7.1 (bm, 6H), 7.22 (d, 2H), 7.35-7.55 (bm, 6H), 7.62 (t, 1H), 7.9 (d, 1H), 8.15 (d, 1H).

15 c) Methyl 2-[(4-{3-[4-(benzyloxy)phenoxy]propyl}phenoxy)methyl]benzoate (0.70 g, 1.45 mmol) was dissolved in dichloromethane (10 ml). Dimethylsulfide (1.08 g, 17.4 mmol) and boron trifluoride etherate (2.06 g, 14.5 mmol) was added and the mixture was stirred at room temperature for six hours. EtOAc (10 ml) was added and the organic phase was washed with water (3 X 10 ml), dried (MgSO_4) and the solvent was removed by evaporation. The crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried (MgSO_4). After removing the solvent by evaporation 0.328 gram of methyl 2-[(4-[3-(4-hydroxyphenoxy)propyl]phenoxy)methyl]benzoate (yield 57.6%) was obtained.

20 ^1H NMR (500 MHz, CDCl_3): δ 2.05 (m, 2H), 2.75 (t, 2H), 3.9 (m, 5H), 5.5 (s, 2H), 6.65-6.8 (bm, 4H), 6.95 (d, 2H), 7.15 (d, 2H), 7.4 (t, 1H), 7.55 (t, 1H), 7.8 (d, 1H), 8.05 (d, 1H).

25 d) Methyl 2-[(4-[3-(4-hydroxyphenoxy)propyl]phenoxy)methyl]benzoate (0.32 g, 0.81 mmol) was dissolved in dichloromethane (10 ml) and cooled to -20° C. Triethylamine (0.123g, 1.22 mmol) was added drop wise and methanesulfonyl chloride (0.10 g, 0.89

mmol) was added drop wise. The ice bath was removed and the mixture was warmed to room temperature. Diethyl ether (5 ml) was added and the precipitate was filtered off. The organic phase was washed with two portions of brine (5 ml) and dried (MgSO_4). Removing the solvent by evaporation gave 0.37 gram of methyl 2-{{4-(3-[(methylsulfonyl)oxy]-phenoxy}propyl)phenoxy}methyl}benzoate (yield 97.3%).

¹HNMR (500 MHz, CDCl_3): δ 2.1 (m, 2H), 2.75 (t, 2H), 3.15 (s, 3H), 3.9 (m, 5H), 5.5 (s, 2H), 6.9-7.0 (bm, 4H), 7.15 (d, 2H), 7.22 (d, 2H), 7.4 (t, 1H), 7.55 (t, 1H), 7.8 (d, 1H), 8.05 (d, 1H).

e) Methyl 2-{{4-(3-[(methylsulfonyl)oxy]phenoxy}propyl)phenoxy}methyl}benzoate (0.38 g, 0.81 mmol) was dissolved in a mixture of THF/water (7/1, 4 ml) and lithium hydroxide (9.3 mg, 0.39 mmol) was added. The reaction was performed in a single node microwave oven (7 min, 150° C). The reaction mixture was acidified (HCl, 1 M, 1 ml) and the water phase was washed with two portions of EtOAc (2 X 5 ml). The organic phases were combined, dried (MgSO_4) and the solvent was removed by evaporation. The crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried (MgSO_4). Removing the solvent by evaporation gave 88 mg of 2-{{4-(3-[(methylsulfonyl)oxy]phenoxy}propyl)phenoxy}methyl}benzoic acid (yield 23.7%).

¹HNMR (500 MHz, CDCl_3): δ 2.1 (m, 2H), 2.75 (t, 2H), 3.15 (s, 3H), 3.95 (t, 2H), 5.58 (s, 2H), 6.9-7.05 (bm, 4H), 7.15-7.25 (bm, 4H), 7.45 (t, 1H), 7.65 (t, 1H), 7.85 (d, 1H), 8.2 (d, 1H).

Example 20

a) 4-(3-Hydroxypropyl)phenol (1.0 g, 6.57 mmol) and methyl 2-(bromomethyl)benzoate (1.66 g, 7.23 mmol) was dissolved in acetonitrile (10 ml). Potassium carbonate (1.82 g, 13.14 mmol) was added and the mixture was stirred at 60° C for three hours. Polymer supported trisamine (0.3 eqv) was added and the solution was stirred at room temperature overnight. The PS-trisamine was filtered off and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic layer was washed with 3 portions of water (3 X 10 ml). The organic phase was dried (MgSO_4) and the solvent was removed by evaporation to give 1.66 gram of methyl 2-[(4-(3-hydroxypropyl)phenoxy)-methyl]benzoate (yield 84.2%).

¹HNMR (500 MHz, CDCl_3): δ 1.9 (m, 2H), 2.65 (t, 2H), 3.25 (s, 1H), 3.65 (t, 2H), 3.85 (s, 3H), 5.45 (s, 2H), 6.95 (d, 2H), 7.15 (d, 2H), 7.35 (t, 1H), 7.5 (t, 1H), 7.75 (d, 1H), 8.05 (d, 1H).

b) Methyl 2-[(4-(3-hydroxypropyl)phenoxy)methyl]benzoate (0.50 g, 1.66 mmol) and 4-(benzyloxy)phenol (0.33 g, 1.66 mmol) was added to a dry round bottomed flask and fitted with a septum. Dry toluene (10 ml) was added and N_2 was flushed through the mixture for 5 minutes. (Tributylphosphoranylidene)acetonitrile (0.80 g, 3.33 mmol) was added drop wise and the reaction was performed in a single node microwave oven. After removing the solvent by evaporation the crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried (MgSO_4). Removing the solvent by evaporation gave 0.515 gram of methyl 2-[(4-(3-[4-(benzyloxy)phenoxy)-propyl]phenoxy)methyl]benzoate (yield 64.1%).

¹HNMR (500 MHz, CDCl₃): δ 2.15 (m, 2H), 2.85 (t, 2H), 4.0 (m, 5H), 5.1 (s, 2H), 5.6 (s, 2H), 6.9-7.1 (bm, 6H), 7.22 (d, 2H), 7.35-7.55 (bm, 6H), 7.62 (t, 1H), 7.9 (d, 1H), 8.15 (d, 1H).

5 c) Methyl 2-[(4-{3-[4-(benzyloxy)phenoxy]propyl}phenoxy)methyl]benzoate (0.70 g, 1.45 mmol) was dissolved in dichloromethane (10 ml). Dimethylsulfide (1.08 g, 17.4 mmol) and borontrifluoride etherate (2.06 g, 14.5 mmol) was added and the mixture was stirred at room temperature for six hours. EtOAc (10 ml) was added and the organic phase was washed with water (3 X 10 ml), dried (MgSO₄) and the solvent was removed by evaporation. The crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried (MgSO₄). After removing the solvent by evaporation 0.328 gram of methyl 2-((4-[3-(4-hydroxyphenoxy)propyl]phenoxy)methyl)benzoate (yield 57.6%) was obtained.

10 ¹HNMR (500 MHz, CDCl₃): δ 2.05 (m, 2H), 2.75 (t, 2H), 3.9 (m, 5H), 5.5 (s, 2H), 6.65-6.8 (bm, 4H), 6.95 (d, 2H), 7.15 (d, 2H), 7.4 (t, 1H), 7.55 (t, 1H), 7.8 (d, 1H), 8.05 (d, 1H).

15 d) Methyl 2-((4-[3-(4-hydroxyphenoxy)propyl]phenoxy)methyl)benzoate (0.32 g, 0.81 mmol) was dissolved in dichloromethane (10 ml) and cooled to -20° C. Triethylamine (0.123 g, 1.22 mmol) was added drop wise and methanesulfonyl chloride (0.10 g, 0.89 mmol) was added drop wise. The ice bath was removed and the mixture was wormed to room temperature. Diethyl ether (5 ml) was added and the precipitate was filtered off. The organic phase was washed with two portions of brine (5 ml) and dried (MgSO₄). Removing the solvent by evaporation gave 0.37 gram of methyl 2-((4-(3-[(methylsulfonyl)oxy]phenoxy)propyl)phenoxy)methyl)benzoate (yield 97.3%).

¹HNMR (500 MHz, CDCl₃): δ 2.1 (m, 2H), 2.75 (t, 2H), 3.15 (s, 3H), 3.9 (m, 5H), 5.5 (s, 2H), 6.9-7.0 (bm, 4H), 7.15 (d, 2H), 7.22 (d, 2H), 7.4 (t, 1H), 7.55 (t, 1H), 7.8 (d, 1H), 8.05 (d, 1H).

5 e) Methyl 2-{{4-[4-[(methylsulfonyl)oxy]phenoxy]propyl}phenoxy}methyl}benzoate (0.38 g, 0.81 mmol) was dissolved in a mixture of THF/water (7/1, 4 ml) and lithium hydroxide (9.3 mg, 0.39 mmol) was added. The reaction was performed in a single node microwave oven (7 min, 150° C). The reaction mixture was acidified (HCl, 1 M, 1 ml) and the water phase was washed with two portions of EtOAc (2 X 5 ml). The organic phases 10 were combined, dried (MgSO₄) and the solvent was removed by evaporation the crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min). The product containing fractions was pooled and the 15 acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried (MgSO₄). Removing the solvent by evaporation gave 63 mg of 2-{{4-[4-hydroxyphenoxy]propyl}phenoxy}methyl}benzoic acid (yield 20.5%).

20 ¹HNMR (500 MHz, CDCl₃): δ 2.05 (m, 2H), 2.75 (t, 2H), 3.9 (t, 2H), 5.58 (s, 2H), 6.65-6.8 (bm, 4H), 6.95 (d, 2H), 7.15 (d, 2H), 7.4 (t, 1H), 7.65 (t, 1H), 7.85 (d, 1H), 8.2 (d, 1H).

Example 21

25 a) 2-(2-Ethoxyphenyl)ethanamine (0.55 g, 3.33 mmol) and 3-(4-hydroxyphenyl)propanoic acid (0.50 g, 3.00 mmol) was dissolved in dimethyl formamide (5 ml) and cooled to 0° C. N-[(1*H*-1,2,3-benzotriazol-1-yl)oxy](dimethylamino)methylene]-*N*-methylmethanaminium tetrafluoroborate (1.18 g, 3.66 mmol) and diisopropylethylamine (0.90 g, 7.0 mmol) were added and the solution was warmed to room temperature and stirred overnight. EtOAc (15 ml) was added and the organic phase was washed with two portions of sodium 30 hydrogencarbonate (aq, 10 ml). The organic phase was dried (MgSO₄) and EtOAc was

removed by evaporation to give 0.98 gram of *N*-[2-(2-ethoxyphenyl)ethyl]-3-(4-hydroxyphenyl)propanamide (yield 93.9%).

¹HNMR (Rotamers, 500 MHz, CDCl₃): δ 1.42 (t, 3H), 2.42 (t, 2H), 2.8-2.92 (m, 4H), 3.55 (q, 2H), 4.05 (q, 2H), 6.08 (m, 1H), 6.82-6.93 (m, 4H), 6.96-7.1 (m, 3H), 7.22 (t, 1H).

b) *N*-[2-(2-ethoxyphenyl)ethyl]-3-(4-hydroxyphenyl)propanamide (0.35 g, 1.12 mmol) and methyl 2-(bromomethyl)benzoate (0.28 g, 1.23 mmol) was dissolved in acetonitrile (5 ml) and potassium carbonate (324 mg, 2.34 mmol) was added. The mixture was stirred at 60° C for three hours. Polymersupported trisamine (0.3 eqv) was added and stirred overnight. The polymer was filtered off, solvent was removed by evaporation, addition of EtOAc (10 ml) and the organic phase was washed with three portions of water. After drying the crude (MgSO₄) and the solvent was removed by evaporation, the crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried (MgSO₄). Removing the solvent by evaporation gave 26 mg of methyl 2-{{4-(3-{{2-(2-ethoxyphenyl)ethyl}amino}-3-oxopropyl)phenoxy}methyl}benzoate (yield 50.4%).

¹HNMR (Rotamers, 500 MHz, CDCl₃): δ 1.38 (t, 3H), 2.35 (t, 2H), 2.76 (t, 2H), 2.85 (t, 2H), 3.45 (q, 2H), 3.86 (s, 3H), 3.99 (q, 2H), 5.44 (s, 2H), 5.84 (m, 1H), 6.78-6.9 (m, 4H), 6.96-7.03 (m, 3H), 7.15 (t, 1H), 7.32 (t, 1H), 7.5 (t, 1H), 7.72 (d, 1H), 8.0 (d, 1H).

c) Methyl 2-{{4-(3-{{2-(2-ethoxyphenyl)ethyl}amino}-3-oxopropyl)phenoxy}methyl}benzoate (0.26 g, 0.56 mmol) was dissolved in a mixture of THF/water (7/1, 5 ml) and lithium hydroxide (54 mg, 2.25 mmol) was added. The reaction was performed in a single node microwave oven (7 min, 150° C). The reaction mixture was acidified (HCl, 1 M, 1 ml) and the water phase was washed with two portions of EtOAc (2 X 5 ml). The organic phases were combined, dried (MgSO₄) and the solvent was removed by evaporation. The

crude product was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were pooled and the 5 acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried ($MgSO_4$). Removing the solvent by evaporation gave 136 mg of 2-{{4-(3-{{2-(2-ethoxyphenyl)ethyl}amino}-3-oxopropyl)phenoxy}methyl}benzoic acid (yield 53.9%).

10 1H NMR (Rotamers, 500 MHz, $CDCl_3$): δ 1.42 (t, 3H), 2.38 (t, 2H), 2.8 (t, 2H), 2.9 (t, 2H), 3.42 (q, 2H), 3.99 (q, 2H), 5.55 (s, 2H), 6.81-6.95 (m, 4H), 7.05-7.17 (m, 4H), 7.38 (t, 1H), 7.55 (m, 1H), 7.81 (d, 1H), 8.11 (d, 1H)

Example 22

15 a) *N*-Ethyl-*N*-(2-pyridin-2-ylethyl)amine (0.5 g, 3.32 mmol) and 3-(4-hydroxyphenyl)-
propanoic acid (0.50 g, 3.00 mmol) was dissolved in dimethylformamide (5 ml) and cooled
o 0° C. *N*-[(1*H*-1,2,3-benzotriazol-1-yloxy)(dimethylamino)methylene]-*N*-
methylmethanaminium tetrafluoroborate (1.18 g, 3.66 mmol) and diisopropylethylamine
(0.90 g, 7.0 mmol) were added and the solution was warmed to room temperature and
20 stirred overnight. EtOAc (15 ml) was added and the organic phase was washed with two
portions of sodium hydrogencarbonate (aq, 10 ml). The organic phase was dried ($MgSO_4$)
and EtOAc was removed by evaporation to give 0.913 gram of *N*-ethyl-3-(4-
hydroxyphenyl)-*N*-(2-pyridin-2-ylethyl)propanamide (yield 91.9%).

1 1H NMR (Rotamers, 500 MHz, $CDCl_3$): δ 0.97, 1.05 (t, t, 3H), 2.41, 2.52 (t, t, 2H), 2.75-
25 3.0 (m, 4H), 3.09, 3.33 (q, q, 2H), 3.54, 3.61 (t, t, 2H), 6.74-6.82 (m, 2H), 6.93-7.2 (m,
4H), 7.58 (m, 1H), 8.48 (m, 1H).

30 b) *N*-Ethyl-3-(4-hydroxyphenyl)-*N*-(2-pyridin-2-ylethyl)propanamide (0.35 g, 1.17 mmol)
and methyl 2-(bromomethyl)benzoate (0.30 g, 1.29 mmol) were dissolved in acetonitrile (5
ml) and potassium carbonate (324 mg, 2.34 mmol) was added. The mixture was stirred at
60° C for three hours. Polymer supported trisamine (0.3 eqv) was added and stirred
overnight. The polymer was filtered off, solvent was removed by evaporation, EtOAc (10

ml) was added and the organic phase was washed with three portions of water. After drying the organic layer ($MgSO_4$), the solvent was removed by evaporation. The crude was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of 5 acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried ($MgSO_4$). Removing the solvent by evaporation gave 135 mg of methyl 2-[(4-{3-[ethyl(2-pyridin-2-ylethyl)amino]-3-10 oxopropyl}phenoxy)methyl]benzoate (yield 25.8%).

¹HNMR (Rotamers, 600 MHz, $CDCl_3$): δ 0.97, 1.05 (t, t, 3H), 2.41, 2.52 (t, t, 2H), 2.75-3.0 (m, 4H), 3.09, 3.33 (q, q, 2H), 3.54, 3.61 (t, t, 2H), 3.83 (s, 3H), 5.43 (s, 2H), 6.75-6.85 (m, 2H), 6.93-7.2 (m, 4H), 7.3 (t, 1H), 5.42 (m, 2H), 7.7 (d, 1H), 7.97 (d, 1H), 8.48 (m, 1H).

c) Methyl 2-[(4-{3-[ethyl(2-pyridin-2-ylethyl)amino]-3-oxopropyl}phenoxy)methyl]-20 benzoate (0.135 g, 0.30 mmol) was dissolved in a mixture of THF/water (7/1, 5 ml) and lithium hydroxide (29 mg, 1.2 mmol) was added. The reaction was performed in a single node microwave oven (7 min, 150° C). The reaction mixture was acidified (HCl, 1 M, 1 ml) and the water phase was washed with two portions of EtOAc (2 X 5 ml). The organic phases were combined, dried ($MgSO_4$) and the solvent was removed by evaporation. The crude product was purified by preparative HPLC (started with acetonitrile/buffer 60/40 and then the acetonitrile concentration was increased to 100%, the buffer was a mixture of 25 acetonitrile/water 10/90 and ammonium acetate (0.1 M, column KR-100-7-C8, 50 mm X 250 mm, flow 40 ml/min)). The product containing fractions were pooled and the acetonitrile was removed by evaporation. EtOAc (10 ml) was added and the organic phase was washed with two portions of brine and dried ($MgSO_4$). Removing the solvent by evaporation gave 27 mg of 2-[(4-{3-[ethyl(2-pyridin-2-ylethyl)amino]-3-30 oxopropyl}phenoxy)methyl]benzoic acid (yield 20.6%).

¹HNMR (Rotamers, 500 MHz, CDCl₃): δ 1.02, 1.12 (t, t, 3H), 2.48, 2.59 (t, t, 2H), 2.85-3.43 (m, 6H), 3.58, 3.66 (t, t, 2H), 5.51, 5.53 (s, s, 2H), 6.86-6.96 (m, 2H), 7.06-7.33 (m, 4H), 7.4 (t, 1H), 5.56 (t, 1H), 7.64-7.75 (m, 2H), 8.14 (m, 1H), 8.64 (m, 1H)

5 Example 23

a) 1-(2-Bromoethyl)-3-*tert*-butoxybenzene

3-(2-Bromoethyl)phenol (1.349 g, 6.709 mmol) in DCM (7 ml) was cooled under argon to 10 -78 °C. Under stirring, isobutene was bubbled into the mixture until more than 5 ml were added. Trifluoromethanesulphonic acid (50 µl) was dropped in. The mixture was stirred under argon at -78 °C for 4.5 hours. Triethylamine (120 µl) was added. The reaction mixture was allowed up to room temperature and then filtered. The filtrate was evaporated to dryness and petroleum ether (25 ml) was added into the residue. It was then filtered and 15 evaporated. The obtained oil was solved in ethyl acetate, washed with water, dried (sodium sulphate) and evaporated. The residue was dissolved in CDCl₃ and then evaporated. 1.223 g desired product was left, yield 71%.

¹H NMR (500 MHz, CDCl₃): δ 1.42 (s, 9H), 3.17 (t, 2H), 3.59 (t, 2H), 6.92-6.98 (m, 3H) and 7.25 (t, 1H).

20

b) Methyl 2-[(2-(3-*tert*-butoxyphenyl)ethyl]thio}benzoate

1-(2-Bromoethyl)-3-*tert*-butoxybenzene (320 mg, 1.244 mmol) was dissolved in 25 acetonitrile (15 ml). Methyl thiosalicylate (209 mg, 1.244 mmol) was added and then potassium carbonate, anhydrous (258 mg, 1.866 mmol) was added. The mixture was heated to reflux for 3 hours and then evaporated under vacuum to dryness.

Chromatography of the residue on a column (ISOLUTE® SI, 5g/25 ml) using ethyl acetate/heptane (5:95) as eluant gave 421 mg desired product, yield 98%.

¹H NMR (500 MHz, CDCl₃): δ 1.39 (s, 9H), 3.00 (t, 2H), 3.21 (t, 2H), 3.96 (s, 3H), 6.90-30 6.93 (m, 2H), 7.01 (d, 1H), 7.19-7.26 (m, 2H), 7.38 (d, 1H), 7.48 (t, 1H) and 8.00 (d, 1H).

c) Methyl 2-{{2-(3-hydroxyphenyl)ethyl}thio}benzoate

Methyl 2-{{2-(3-*tert*-butoxyphenyl)ethyl}thio}benzoate (402 mg, 1.167 mmol) was dissolved in DCM (3 ml). Trifluoroacetic acid (3 ml) was added. The mixture was stirred overnight and then evaporated to dryness. Chromatography of the residue on a column (ISOLUTE® SI, 2g/6ml) using ethyl acetate/heptane (2.5:97.5, then 5:95, then 10:90 and then 25:75) as eluant gave 260 mg the desired product, yield 77%.

¹H NMR (500 MHz, CDCl₃): δ 2.94 (t, 2H), 3.17 (t, 2H), 3.95 (s, 3H), 6.18 (s, 1H) 6.77-6.83 (m, 3H), 7.18 (t, 2H), 7.34 (d, 1H), 7.44 (t, 1H) and 7.99 (d, 1H).

10

d) N-benzyl-2-bromo-N-hexylacetamide

N-hexylbenzylamine (4.2 g, 21.953 mmol) and triethylamine (3.98 ml, 28.539 mmol) were mixed in DCM (20 ml) and cooled in an ice-bath. Bromoacetyl chloride (3.455 mg, 21.953 mmol) in DCM (5 ml) was added. The mixture was stirred over weekend and the temperature was allowed going up to room temperature. The mixture was washed with water mixed with 1% hydrochloric acid (water phase pH~4-5) and brine, dried with magnesium sulphate, and evaporated. Column chromatography of the residue on silica gel using ethyl acetate/heptane (10:90, then 20:80) as eluant gave 4.0 g desired product, yield 58%.

¹H NMR (rotamers, 300 MHz, CDCl₃): δ 0.85-0.92 (m, 3H), 1.28 (s, br, 6H), 1.53-1.62 (m, 2H), 3.25, 3.39 (t, t, 2H), 4.05, 4.16 (s, s, 2H), 4.61, 4.63 (s, s, 2H) and 7.19 -7.42 (m, 5H).

25

e) Methyl 2-{{2-(3-{2-[benzyl(hexyl)amino]-2-oxoethoxy}phenyl)ethyl}thio}benzoate

Methyl 2-{{2-(3-hydroxyphenyl)ethyl}thio}benzoate (129 mg, 0.447 mmol), *N*-benzyl-2-bromo-*N*-hexylacetamide (154 mg, 0.492 mmol) and potassium carbonate, anhydrous (93 mg, 0.671 mmol) were mixed in acetonitrile (10 ml). The mixture was heated to reflux overnight and then evaporated to dryness. Water and ethyl acetate were added into the residue. The two phases were separated. The organic phase was washed with brine and dried with magnesium sulphate and then evaporated. Chromatography of the residue on a

column (ISOLUTE® SI, 2g/6ml) using ethyl acetate/heptane (10:90, then 25:75) as eluant gave 208 mg the desired product, yield 89.5%.

¹H NMR (rotamers, 500 MHz, CDCl₃): δ 0.86-0.91 (m, 3H), 1.24-1.32 (m, 6H), 1.53-1.64 (m, 2H), 2.94-3.03 (m, 2H), 3.13-3.20 (m, 2H), 3.29, 3.41 (t, t, 2H), 3.93 (s, 3H), 4.64, 4.65 (s, s, 2H), 4.71, 4.81 (s, s, 2H), 6.75-6.77 (m, 1H), 6.85-6.93 (m, 2H), 7.17-7.39 (m, 8H), 7.46 (t, 1H) and 7.99 (d, 1H).

¹³C NMR (rotamers, 125 MHz, CDCl₃): δ 13.85, 13.86, 22.39, 26.38, 27.03, 28.27, 31.31, 31.37, 33.28, 34.51, 34.57, 46.34, 48.13, 50.33, 51.94, 67.13, 67.38, 112.33, 112.39, 114.84, 114.96, 121.48, 121.56, 123.79, 125.49, 126.40, 127.24, 127.54, 127.64, 127.89, 128.42, 128.74, 129.48, 129.57, 131.17, 132.22, 136.51, 137.03, 141.30, 141.74, 141.87, 158.06, 158.15, 166.75, 167.74 and 167.88.

f) 2-{{2-(3-{2-[Benzyl(hexyl)amino]-2-oxoethoxy}phenyl)ethyl}thio}benzoic acid

Methyl 2-{{2-(3-{2-[benzyl(hexyl)amino]-2-oxoethoxy}phenyl)ethyl}thio}benzoate (80 mg, 0.154 mmol) in tetrahydrofuran (3 ml) was cooled in an ice-bath. Lithium hydroxide (7.4 mg, 0.308 mmol) in water (3 ml) was added. The cooling bath was then removed and the mixture was stirred for 12 days and then evaporated in vacuum to remove tetrahydrofuran. The residue was acidified with 1% hydrochloric acid, pH=3, and extracted with ethyl acetate. The organic phase was dried with magnesium sulphate and evaporated. Chromatography of the residue on a column (ISOLUTE® SI, 2g/6 ml) using DCM, MeOH/DCM (1:99, and then 2:98) as eluant gave 27 mg product mixture. Re-chromatography of the mixture on a column (ISOLUTE® SI, 1g/6ml) using DCM and then MeOH/AcOH/DCM (0.25:0.25:99.5) as eluant gave 17 mg desired product, yield 22%.

¹H NMR (rotamers, 500 MHz, CDCl₃): δ 0.85-0.90 (m, 3H), 1.23-1.31 (m, 6H), 1.53-1.63 (m, 2H), 2.91-3.00 (m, 2H), 3.11-3.20 (m, 2H), 3.29, 3.41 (t, t, 2H), 4.65, 4.66 (s, s, 2H), 4.74, 4.83 (s, s, 2H), 6.72-6.93 (m, 3H), 7.20-7.32 (m, 6H), 7.37 (t, 2H), 7.47 (t, 1H) and 8.09 (d, 1H).

¹³C NMR (rotamers, 125 MHz, CDCl₃): δ 13.98, 22.51, 26.51, 27.11, 28.36, 31.42, 31.49, 34.16, 34.60, 34.66, 46.52, 48.35, 50.52, 67.20, 67.47, 112.60, 115.15, 115.29, 121.66,

121.75, 124.46, 126.54, 126.80, 127.42, 127.70, 128.04, 128.56, 128.89, 129.59, 129.67,
132.13, 132.79, 136.46, 136.98, 141.16, 141.75, 141.89, 158.08, 158.17, 168.25, 168.37
and 169.60.

5 **Biological activity**

Formulations

Compounds were dissolved in DMSO to obtain 16 mM stock solutions. Before assays,
stock solutions were further diluted in DMSO and culture media.

10

GENERAL CHEMICALS AND REAGENTS

Luciferase assay reagent was purchased from Packard, USA. Restriction Enzymes were
from Boehringer and Vent polymerase from New England Biolabs.

15 **CELL LINES AND CELL CULTURE CONDITIONS**

U2-OS, (Osteogenic sarcoma, Human) was purchased from ATCC, USA. Cells were
expanded and refrozen in batches from passage number six. Cells were cultured in
Dulbecco's modified Eagle medium (DMEM) with 25 mM glucose, 2 mM glutamine or 4
mM L-alanyl-L-glutamine, 10% fetal calf serum, at 5% CO₂. Phosphate buffered saline
20 (PBS) without addition of calcium or magnesium was used. All cell culture reagents were
from Gibco (USA) and 96-well cell culture plates were purchased from Wallach.

PLASMID CONSTRUCTS FOR HETEROLOGOUS EXPRESSION

Standard recombinant DNA techniques were carried out as described by Ausubel (7). The
25 Luciferase reporter vector, pGL5UAS (clone consists of five copies of the GAL4 DNA
binding sequence, 5'-CGACGGAGTACTGTCCTCCGAGCT-3', cloned into the
SacI/XhoI sites of pGL3-Promoter (Promega). The SacI/XhoI fragment carrying the UAS
sites was constructed using annealed overlapping oligonucleotides.

Expression vectors used are based upon pSG5 (Stratagene). All vectors contain an EcoRI/NheI fragment encoding the DNA binding domain of GAL4 (encoding amino acid positions 1-145 of database accession number P04386) followed by an in-frame fusion to a fragment encoding the nuclear localisation sequence from T antigen of Polyoma Virus.

5 The nuclear localisation sequence was constructed using annealed overlapping oligonucleotides creating NheI/KpnI sticky ends (5'-CTAGCGCTCCTAGAAGAAACGCAAGGTTGGTAC-3'). The ligand binding domains from human and mouse PPAR α and human and mouse PPAR γ were PCR amplified as KpnI/BamHI fragments and cloned in frame to the GAL4 DNA binding domain and the nuclear localisation sequence. The sequence of all plasmid constructs used were confirmed by sequencing.

10

The following expression vectors were used for transient transfections:

vector	encoded PPAR subtype	sequence reference ¹
pSGGALhPPa	human PPAR α	S74349, nt 625-1530
pSGGALmPPa	murine PPAR α	X57638, nt 668-1573
pSGGALhPPg	human PPAR γ	U63415, nt 613-1518
pSGGALmPPg	murine PPAR γ	U09138, nt 652-1577

15 ¹ refers to nucleotide positions of data base entry used to express the ligand binding domain.

TRANSIENT TRANSFECTIONS

Frozen stocks of cells from passage number six were thawed and expanded to passage number eight before transfections. Confluent cells were trypsinised, washed and pelleted by centrifugation at 270xg for 2 minutes. The cell pellet was resuspended in cold PBS to a cell concentration of about 18×10^6 cells/ml. After addition of DNA, the cell suspension

was incubated on ice for approximately 5 minutes before electroporation at 230 V, 960 μ F in Biorad's Gene PulserTM in 0.5 ml batches. A total of 50 μ g DNA was added to each batch of 0.5 ml cells, including 2.5 μ g expression vector, 25 μ g reporter vector and 22.5 μ g unspecific DNA (pBluescript, Stratagene).

5

After electroporation, cells were diluted to a concentration of 320'000 cells/ml in DMEM without phenol red, and approximately 25'000 cells/well were seeded in 96-well plates. In order to allow cells to recover, seeded plates were incubated at 37°C for 3-4 hours before addition of test compounds. In assays for PPAR α , the cell medium was supplemented with 10 resin-charcoal stripped fetal calf serum (FCS) in order to avoid background activation by fatty acid components of the FCS. The resin-charcoal stripped FCS was produced as follows; for 500 ml of heat-inactivated FCS, 10 g charcoal and 25 g Bio-Rad Analytical Grade Anion Exchange Resin 200-400 mesh were added, and the solution was kept on a magnetic stirrer at room temperature over night. The following day, the FCS was 15 centrifuged and the stripping procedure was repeated for 4-6 hours. After the second treatment, the FCS was centrifuged and filter sterilised in order to remove remnants of charcoal and resin.

ASSAY PROCEDURE

20 Stock solutions of compounds in DMSO were diluted in appropriate concentration ranges in master plates. From master plates, compounds were diluted in culture media to obtain test compound solutions for final doses.

25 After adjustment of the amount of cell medium to 75 μ l in each well, 50 μ l test compound solution was added. Transiently transfected cells were exposed to compounds for about 24 hours before the luciferase detection assay was performed. For luciferase assays, 100 μ l of assay reagent was added manually to each well and plates were left for approximately 20 minutes in order to allow lysis of the cells. After lysis, luciferase activity was measured in a 1420 Multiwell counter, Victor, from Wallach.

Reference compounds

The TZD pioglitazone was used as reference substance for activation of both human and murine PPAR γ . 5,8,11,14-Eicosatetrayonic acid (ETYA) was used as reference substance for human PPAR α .

Calculations and analysis

For calculation of ED₅₀ values, a concentration-effect curve was established. Values used were derived from the average of two or three independent measurements (after subtraction of the background average value) and were expressed as the percentage of the maximal activation obtained by the reference compound. Values were plotted against the logarithm of the test compound concentration. ED₅₀ values were estimated by linear intercalation between the data points and calculating the concentration required to achieve 50% of the maximal activation obtained by the reference compound.

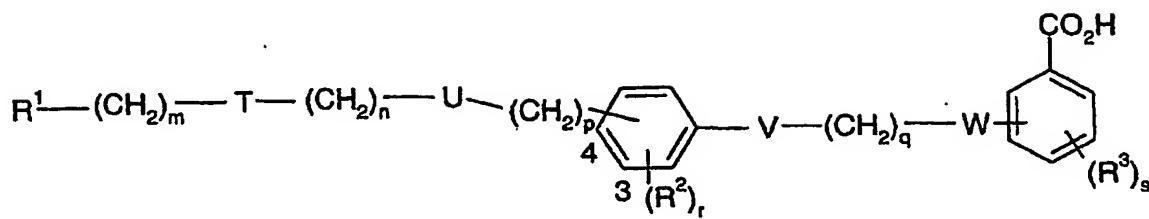
15

The compound of formula I have an ED₅₀ of less than 50 μ mol for PPAR α and/or PPAR γ . Preferred compounds have an ED₅₀ of less than 5 μ mol for either PPAR α or PPAR γ .

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CLAIMS

1. A compound of formula I



5 wherein

R¹ represents aryl optionally substituted by a heterocyclic group or a heterocyclic group optionally substituted by aryl wherein each aryl or heterocyclic group is optionally substituted by one or more of the following groups:

- 10 a C₁₋₆alkyl group;
- a C₁₋₆acyl group;
- arylC₁₋₆alkyl, wherein the alkyl, aryl, or alkylaryl group is optionally substituted by one or more R^b;
- halogen,
- 15 -CN and NO₂,
- NR^cCOOR^a;
- NR^cCOR^a;
- NR^cR^a;
- NR^cSO₂R^d;
- 20 -NR^cCONR^kR^c;
- NR^cCSNR^aR^k;
- OR^a;
- OSO₂R^d;
- SO₂R^d;
- 25 -SOR^d;
- SR^c;
- SO₂NR^aR^f;

-SO₂OR^a;

-CONR^cR^a;

-OCONR^fR^a;

wherein R^a represents H, a C₁₋₆alkyl group, aryl or arylC₁₋₆alkyl group wherein the alkyl,

5 aryl or arylC₁₋₆alkyl group is optionally substituted one or more times by R^b, wherein R^b represents C₁₋₆alkyl, aryl, arylC₁₋₆alkyl, cyano, -NR^cR^d, =O, halo, -OH, -SH, -OC₁₋₄alkyl, -Oaryl, -OC₁₋₄alkylaryl, -COR^c, -SR^d, -SOR^d, or -SO₂R^d, wherein R^c represents H, C₁₋₄alkyl, aryl, arylC₁₋₄alkyl and R^d represents C₁₋₄alkyl, aryl, arylC₁₋₄alkyl;

10 wherein R^f represents hydrogen, C₁₋₄alkyl, C₁₋₄acyl, aryl, arylC₁₋₄alkyl and R^a is as defined above; and

R^k represents hydrogen, C₁₋₄alkyl, aryl, arylC₁₋₄alkyl;

m represents 0, 1, 2, 3, 4 or 5 ;

15 T represents O, S, NC(O)N(R⁴), S(O₂)N(R⁵), (R⁵)NS(O₂), N(R⁶)C(O), C(O)N(R⁷), or a single bond;

n represents 0, 1, 2, 3, 4 or 5 ;

20 U represents O, S or a single bond provided that when U is O or S then n represents 1, 2, 3, 4 or 5 and further provided that T and U do not both represent a single bond simultaneously;

p represents 0, 1, 2, 3, 4 or 5 ;

25 wherein the group (CH₂)_p, or the group U if p is 0, is attached at the 3 or 4 position of the phenyl ring as indicated in formula I

V represents O, S, NR⁸, or a single bond;

30 q represents 1, 2 or 3 ;

W represents O, S, N(R⁹)C(O), NR¹⁰, or a single bond;

R² represents halo, a C₁₋₄alkyl group which is optionally substituted by one or more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro, a C₁₋₄acyl group, 5 aryl, an arylC₁₋₄alkyl group, CN or NO₂;

r represents 0, 1, 2 or 3;

R³ represents represents halo, a C₁₋₄alkyl group which is optionally substituted by one or 10 more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro, a C₁₋₄acyl group, aryl, an arylC₁₋₄alkyl group, or CN;

s represents 0, 1, 2 or 3; and

15 R⁴, R⁵, R⁶, R⁷, R⁸, R⁹ and R¹⁰ independently represent H, a C₁₋₁₀alkyl group, aryl or an arylC₁₋₄alkyl group or when m is 0 and T represents a group N(R⁶)C(O) or a group (R⁵)NS(O₂) then R¹ and R⁶ or R¹ and R⁵ together with the nitrogen atom to which they are attached represent a heteroaryl group; as well as optical isomers and racemates thereof as well as pharmaceutically acceptable 20 salts, prodrugs, solvates and crystalline forms thereof.

2. A compound according to claim 1 in which R¹ represents phenyl which is optionally substituted by one or more of the following: halo, hydroxy, a C₁₋₄alkyl group which is 25 optionally substituted by one or more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro, benzyloxy, a C₁₋₄alkylsulphonyloxy group, phenyl or a heteroaryl group, or R¹ represents heteroaryl which is optionally substituted by one or more of the following: halo, a C₁₋₄alkyl group which is optionally substituted by one or more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro or phenyl optionally substituted by one or more of the following: halo, a C₁₋₄alkyl group 30 which is optionally substituted by one or more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro.

3. A compound according to either claim 1 or claim 2 in which the group $-(CH_2)_m-T-(CH_2)_n-U-(CH_2)_p-$ is attached at either the 3 or 4 position in the phenyl ring as indicated by the numbers in formula I and represents a group selected from: $O(CH_2)_2$, $O(CH_2)_3$, $NC(O)NR^4(CH_2)_2$, $CH_2S(O_2)NR^5(CH_2)_2$, $CH_2N(R^6)C(O)CH_2$, $(CH_2)_2N(R^6)C(O)(CH_2)_2$,
5 $C(O)NR^7CH_2$, $C(O)NR^7(CH_2)_2$, $CO(CH_2)_2$, and $CH_2N(R^6)C(O)CH_2O$, wherein R^4 , R^5 ,
 R^6 , and R^7 are as previously defined.

4. A compound according to any previous claim in which the group $-(CH_2)_m-T-(CH_2)_n-U-(CH_2)_p-$ is attached at the 4 position in the phenyl ring as indicated by the numbers in
10 formula I, that is para to the group V.

5. A compound according to any previous claim in which the group $-V-(CH_2)_q-W-$ represents a group selected from : OCH_2 , SCH_2 , $NHCH_2$, CH_2CH_2S or CH_2CH_2O .

15 6. A compound according to any previous claim in which the group $-V-(CH_2)_q-W-$ represents the group OCH_2 .

7. A compound according to any previous claim in which the group $-V-(CH_2)_q-W-$ is joined at the ortho position with respect to the carboxylic acid group.

20 8. A compound according to any previous claim in which R^2 is halo, a C_{1-4} alkyl group or a C_{1-4} alkoxy group and r is 0 or 1.

9. A compound selected from

25 3-{{(3-{{(1,1'-biphenyl-4-ylcarbonyl)amino]methyl}phenyl)amino]methyl}benzoic acid;
2-{{4-(2-oxo-2-{{4-(trifluoromethyl)benzyl]amino}ethyl)phenoxy]methyl}benzoic acid;
2-[(3-{2-[benzyl(hexyl)amino]-2-oxoethyl}phenoxy)methyl]benzoic acid;
2-[(3-(2-oxo-2-{{4-(trifluoromethyl)benzyl]amino}ethyl)phenoxy)methyl]benzoic acid;

2-[(4-{3-[[2-(3,4-dimethoxyphenyl)ethyl](methyl)amino]-3-oxopropyl}phenoxy)-methyl]benzoic acid;

2-[(4-{2-[(4-methyl-2-[4-(trifluoromethyl)phenyl]-1,3-thiazol-5-yl]carbonyl)amino}-ethyl}phenoxy)methyl]benzoic acid;

5 2-({4-[2-((2,4-difluorophenyl)amino]carbonyl)amino}ethyl]phenoxy)methyl]benzoic acid;

2-[(4-{2-[(2-methyl-5-phenyl-3-furoyl)amino]ethyl}phenoxy)methyl]benzoic acid;

2-[(4-{2-[(benzylsulfonyl)amino]ethyl}phenoxy)methyl]benzoic acid;

2-[(4-{2-[benzyl(hexyl)amino]-2-oxoethyl}-2-fluorophenoxy)methyl]benzoic acid;

10 2-[(4-{2-[benzyl(hexyl)amino]-2-oxoethyl}-2-methoxyphenoxy)methyl]benzoic acid;

2-({4-[3-(3,4-dihydroisoquinolin-2(1H)-yl)-3-oxopropyl]phenoxy)methyl]benzoic acid;

2-[(4-{2-[4-(1H-imidazol-1-yl)phenoxy]ethyl}-phenoxy)methyl]benzoic acid;

2-{{4-(2-{4-[(methylsulfonyl)oxy]phenoxy}ethyl)phenoxy)methyl]benzoic acid;

2-[(3-{2-[4-(benzyloxy)phenoxy]ethyl}phenoxy)methyl]benzoic acid;

15 2-{{3-(2-{4-[(methylsulfonyl)oxy]phenoxy}ethyl)phenoxy)methyl]benzoic acid;

2-{{3-[2-(4-hydroxyphenoxy)ethyl]phenoxy)methyl]benzoic acid;

2-[(4-{3-[4-(benzyloxy)phenoxy]propyl}phenoxy)methyl]benzoic acid;

2-{{4-(3-{4-[(methylsulfonyl)oxy]phenoxy}propyl)phenoxy)methyl]benzoic acid;

2-{{4-[3-(4-hydroxyphenoxy)propyl]phenoxy)methyl]benzoic acid;

20 2-{{4-(3-{2-(2-ethoxyphenyl)ethyl}amino)-3-oxopropyl}phenoxy)methyl]benzoic acid;

2-[(4-{3-[ethyl(2-pyridin-2-ylethyl)amino]-3-oxopropyl}phenoxy)methyl]benzoic acid;

and 2-{{2-(3-{2-[benzyl(hexyl)amino]-2-oxoethoxy}phenyl)ethyl}thio}benzoic acid;

and pharmaceutically acceptable salts, prodrugs, solvates and crystalline forms thereof.

25 10. A pharmaceutical formulation comprising a compound according to any preceding claim in admixture with pharmaceutically acceptable adjuvants, diluents and/or carriers.

11. A method of treating or preventing insulin resistance comprising the administration of a compound according to any one of claims 1 to 9 to a mammal in need thereof.

30 12. The use of a compound according to any one of claims 1 to 9 in the manufacture of a medicament for the treatment of insulin resistance.

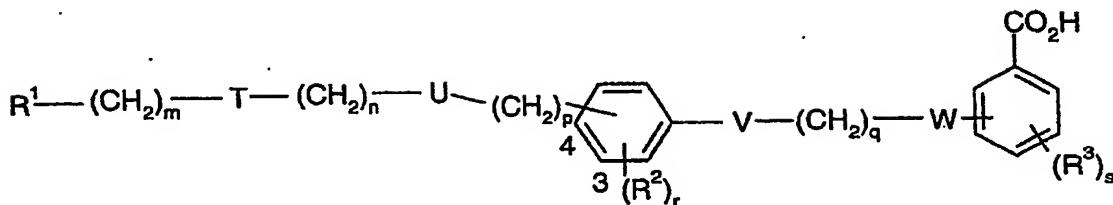
13. Processes to prepare compounds of formula I as described herein.

14. Compounds of formula II.

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ABSTRACT

A compound of formula I



5 wherein

- R¹ represents aryl optionally substituted by a heterocyclic group or a heterocyclic group optionally substituted by aryl wherein each aryl or heterocyclic group is optionally substituted by one or more of the following groups:
 - a C₁-₆alkyl group;
 - 10 a C₁-₆acyl group;
 - arylC₁-₆alkyl, wherein the alkyl, aryl, or alkylaryl group is optionally substituted by one or more R^b;
 - halogen,
 - CN and NO₂,
 - 15 -NR^cCOOR^a;
 - NR^cCOR^a;
 - NR^cR^a;
 - NR^cSO₂R^d;
 - NR^cCONR^kR^c;
 - 20 -NR^cCSNR^aR^k;
 - OR^a;
 - OSO₂R^d;
 - SO₂R^d;
 - SOR^d;
 - 25 -SR^c;
 - SO₂NR^aR^f;
 - SO₂OR^a;

-CONR^cR^a;

-OCONR^fR^a;

wherein R^a represents H, a C₁₋₆alkyl group, aryl or arylC₁₋₆alkyl group wherein the alkyl,

aryl or arylC₁₋₆alkyl group is optionally substituted one or more times by R^b, wherein R^b

5 represents C₁₋₆alkyl, aryl, arylC₁₋₆alkyl, cyano, -NR^cR^d, =O, halo, -OH, -SH, -OC₁₋₄alkyl, -

Oaryl, -OC₁₋₄alkylaryl, -COR^c, -SR^d, -SOR^d, or -SO₂R^d, wherein R^c represents H, C₁₋

4alkyl, aryl, arylC₁₋₄alkyl and R^d represents C₁₋₄alkyl, aryl, arylC₁₋₄alkyl;

wherein R^f represents hydrogen, C₁₋₄alkyl, C₁₋₄acyl, aryl, arylC₁₋₄alkyl and R^a is as defined above; and

10 R^k represents hydrogen, C₁₋₄alkyl, aryl, arylC₁₋₄alkyl;

m represents 0, 1, 2, 3, 4 or 5 ;

T represents O, S, NC(O)N(R⁴), S(O₂)N(R⁵), (R⁵)NS(O₂), N(R⁶)C(O), C(O)N(R⁷), or a

15 single bond;

n represents 0, 1, 2, 3, 4 or 5 ;

U represents O, S or a single bond provided that when U is O or S then n represents 1, 2, 3,

20 4 or 5 and further provided that T and U do not both represent a single bond

simultaneously;

p represents 0, 1, 2, 3, 4 or 5 ;

25 wherein the group (CH₂)_p, or the group U if p is 0, is attached at the 3 or 4 position of the phenyl ring as indicated in formula I

V represents O, S, NR⁸, or a single bond;

30 q represents 1, 2 or 3 ;

W represents O, S, N(R⁹)C(O), NR¹⁰, or a single bond;

R² represents halo, a C₁₋₄alkyl group which is optionally substituted by one or more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro, a C₁₋₄acyl group, aryl, an arylC₁₋₄alkyl group, CN or NO₂;

5

r represents 0, 1, 2 or 3;

R³ represents represents halo, a C₁₋₄alkyl group which is optionally substituted by one or more fluoro, a C₁₋₄alkoxy group which is optionally substituted by one or more fluoro, a C₁₋₄acyl group, aryl, an arylC₁₋₄alkyl group, or CN;

s represents 0, 1, 2 or 3; and

R⁴, R⁵, R⁶, R⁷, R⁸, R⁹ and R¹⁰ independently represent H, a C₁₋₁₀alkyl group, aryl or an arylC₁₋₄alkyl group or when m is 0 and T represents a group N(R⁶)C(O) or a group (R⁵)NS(O₂) then R¹ and R⁶ or R¹and R⁵ together with the nitrogen atom to which they are attached represent a heteroaryl group; as well as optical isomers and racemates thereof as well as pharmaceutically acceptable salts, prodrugs, solvates and crystalline forms thereof processes for preparing such compounds, their utility in treating clinical conditions associated with insulin resistance, methods for their therapeutic use and pharmaceutical compositions containing them.

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